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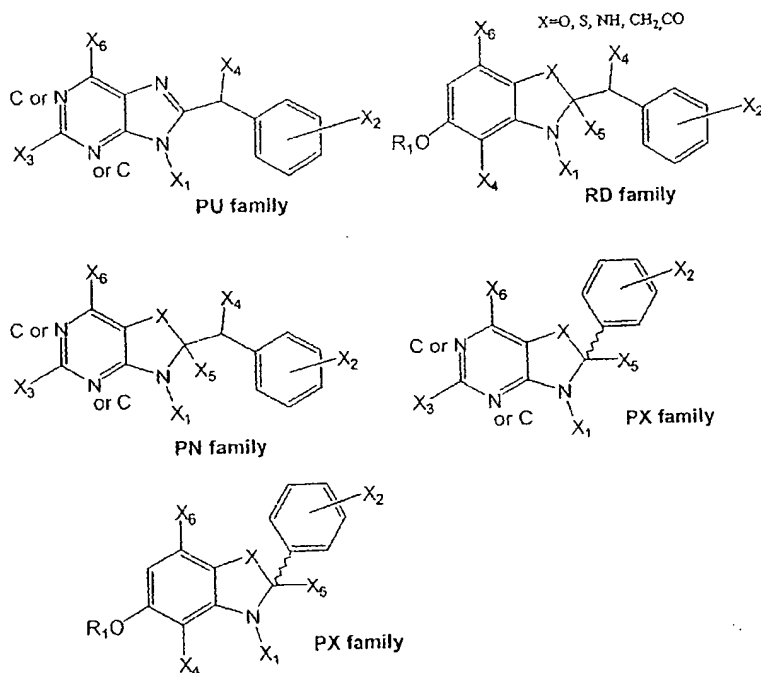
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(54) Title: SMALL MOLECULE COMPOSITIONS FOR BINDING TO HSP90



(57) Abstract: Structural differences in binding pockets of members of the HSP90 family can be exploited to achieve differential degradation of kinases and other signaling proteins through the use of designed small molecules which interact with the N-terminal binding pocket with an affinity which is greater than ADP and different from the ansamycin antibiotics for at least one species of the HSP90 family. Moreover, these small molecules can be designed to be soluble in aqueous media, thus providing a further advantage over the use of ansamycin antibiotics. Pharmaceutical compositions can be formulated containing a pharmaceutically acceptable carrier and a molecule that includes a binding moiety which binds to the N-terminal pocket of at least one member of the HSP90 family of proteins. Such binding moieties were found to have antiproliferative activity against tumor cells which are dependent on proteins requiring chaperones of the HSP90

family for their function. Different chemical species have different activity, however, allowing the selection of, for example Her2 degradation without degradation of Raf kinase. Thus, the binding moieties possess an inherent targeting capacity. In addition, the small molecules can be linked to targeting moieties to provide targeting of the activity to specific classes of cells. Thus, the invention further provides a method for treatment of diseases, including cancers, by administration of these compositions. Dimeric forms of the binding moieties may also be employed.



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SMALL MOLECULE COMPOSITIONS FOR BINDING TO HSP90

This application claims the benefit of US Provisional Application No. 60/245,177, filed November 2, 2000, which is incorporated herein by reference.

Background of the Invention

This application relates to small molecules that bind to the HSP90 family of proteins, and to methods for making and using such small molecules.

The HSP90 family of proteins has four recognized members in mammalian cells: Hsp90 α and β , Grp94 and Trap-1. Hsp90 α and β exist in the cytosol and the nucleus in association with a number of other proteins. Hsp90 is the most abundant cellular chaperone, and has been shown in experimental systems to be required for ATP-dependent refolding of denatured or "unfolded" proteins. It has therefore been proposed to function as part of the cellular defense against stress. When cells are exposed to heat or other environmental stresses, the aggregation of unfolded proteins is prevented by pathways that catalyze their refolding or degradation. This process depends on the association of the unfolded protein in an ordered fashion with multiple chaperones (Hsp 60, 90 and 70 and p23), forming a "refoldosome" and ultimately the ATP-dependent release of the chaperones from the refolded protein.

Hsp90 may also play a role in maintaining the stability and function of mutated proteins. It seems to be required for expression of mutated p53 and v-src to a much greater extent than for their wild-type counterparts. It has been suggested that this occurs as a result of hsp90-mediated suppression of the phenotypes of mutations that lead to protein unfolding.

Hsp90 is also necessary to the conformational maturation of several key proteins involved in the growth response of the cell to extracellular factors. These include the steroid receptors as well as certain transmembrane kinases (i.e., Raf serine kinase, v-src and Her2). The mechanism whereby hsp90 affects these proteins is not fully understood, but appears to be similar to its role in protein refolding. In the case of the progesterone receptor, it has been shown that binding and release of hsp90 from the receptor occurs in a cyclic fashion in concert with release of other chaperones and

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immunophilins and is required for high affinity binding of the steroid to the receptor. Thus, hsp90 could function as a physiologic regulator of signaling pathways, even in the absence of stress.

Hsp90 has also been shown to be overexpressed in multiple tumor types and as a function of oncogenic transformation. Whether it plays a necessary role in maintaining transformation is unknown, but it could have at least three functions in this regard. Cancer cells grow in an environment of hypoxia, low pH and low nutrient concentration. They also rapidly adapt to or are selected to become resistant to radiation and cytotoxic chemotherapeutic agents. Thus, the general role of hsp90 in maintaining the stability of proteins under stress may be necessary for cell viability under these conditions. Secondly, cancer cells harbor mutated oncogenic proteins. Some of these are gain-of-function mutations which are necessary for the transformed phenotype. Hsp90 may be required for maintaining the folded, functionally-active conformation of these proteins. Thirdly, activation of signaling pathways mediated by steroid receptors, Raf and other hsp90 targets is necessary for the growth and survival of many tumors which thus probably also require functional hsp90.

These characteristics of hsp90 make it a viable target for therapeutic agents. HSP90 family members possess a unique pocket in their N-terminal region that is specific to and conserved among all hsp90s from bacteria to mammals, but which is not present in other molecular chaperones. The endogenous ligand for this pocket is not known, but it binds ATP and ADP with low affinity and has weak ATPase activity. In addition, the ansamycin antibiotics geldanamycin (GM) and herbimycin (HA) have been shown to bind to this conserved pocket. This binding affinity has been shown for all members of the HSP90 family. International Patent Publication No. WO98/51702, which is incorporated herein by reference, discloses the use of ansamycin antibiotics coupled to a targeting moiety to provide targeted delivery of the ansamycin leading to the degradation of proteins in and death of the targeted cells. International Patent Publication No. WO00/61578 relates to bifunctional molecules having two moieties which interact with the chaperone protein hsp90, including in particular homo- and heterodimers of ansamycin antibiotics. These bifunctional molecules act to promote degradation and/or

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inhibition of HER-family tyrosine kinases and are effective for treatment of cancers which overexpress Her-kinases.

While the use of geldanamycin and other ansamycin antibiotics and their derivatives provides for effective degradation of a number of kinases and other signaling proteins, they generally lack significant selectivity, and are instead effective to degrade a broad spectrum of proteins. This can increase the risk of undesirable side effects. Furthermore, ansamycin antibiotics are insoluble or at best poorly soluble in aqueous media. This complicates administration. Thus, there remains room for improvement of therapeutic agents that bring about the degradation of kinases and other signaling proteins via interaction with members of the HSP90 family of chaperone proteins.

Summary of the Invention

While the members of the HSP90 family of proteins are characterized by a unique N-terminal binding pocket that is highly conserved, there are structural differences between the pockets of the various members. It has now been found that these structural differences can be exploited to achieve differential degradation of kinases and other signaling proteins through the use of designed small molecules which interact with the N-terminal binding pocket with an affinity which is greater than ADP and different from the ansamycin antibiotics. Moreover, these small molecules can be designed to be soluble in aqueous media, thus providing a further advantage over the use of ansamycin antibiotics.

The N-terminal pockets of the HSP90 of family of proteins contain five potential binding sites. In the case of hsp90 α , these binding sites are:

- (a) a top binding site comprising Lys112,
- (b) a second top binding site comprising Lys58, Asp54, Phe138 backbone, Gly 135 and Asn106;
- (c) a bottom binding site comprising Asp93, Ser52 and Asn51;
- (d) a hydrophobic lateral binding site comprising Val150, Met98, Val186, Phe138, Leu107, Leu103, Val186 and Trp162; and
- (e) a small hydrophobic bottom binding site comprising Thr184, Val186, Val150 and Ile91.

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The present invention provides pharmaceutical compositions comprising a pharmaceutically acceptable carrier and a molecule comprising a binding moiety which binds to the N-terminal pocket of at least one member of the HSP90 family of proteins. This binding moiety interacts with the N-terminal pocket with an affinity greater than ADP but less than geldanamycin for at least one specific protein in the HSP90 family. Further, the binding moiety has a backbone which can achieve a folded C-configuration when disposed within the N-terminal pocket of a member of the HSP90 family of proteins. The binding moiety also has substituents on the backbone directed in orientations to interact with a plurality of the potential binding sites within the N-terminal pocket.

The binding moieties of the invention were found to have antiproliferative activity against tumor cells which are dependent on proteins requiring chaperones of the HSP90 family for their function. Different chemical species have different activity, however, allowing the selection of, for example Her2 degradation without degradation of Raf kinase. Thus, the binding moieties of the invention possess an inherent targeting capacity. In addition, the small molecules can be linked to targeting moieties to provide targeting of the activity to specific classes of cells. Thus, the invention further provides a method for treatment of diseases, including cancers, by administration of these compositions. Dimeric forms of the binding moieties may also be employed.

Brief Description of the Drawings

Fig. 1 shows aligned sequences of amino acids contributing to the N-terminal binding pocket in known members of the HSP90 family of chaperone proteins, Hsp90 α (Seq. ID No. 1), GRP94 (Seq. ID. No. 2), Hsp90 β (Seq. ID No. 3) and Trap1 (Seq. ID. No. 4);

Fig. 2 shows a three dimensional drawings of the pocket with a binding moiety of the invention disposed therein;

Figs. 3A and B show the conformation of PU3 inside the hsp90 pocket, as determined by molecular modeling, and outside of the pocket as determined by x-ray crystallography;

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Fig. 4 shows an exemplary binding moiety/compound based on a purine nucleus, and indicates the interactions of the various substituents with binding sites in the N-terminal pocket of members of the HSP90 family;

Fig. 5 shows the structure of compounds in accordance with the invention;

Fig. 6A shows a synthetic procedure for making compounds in accordance with the invention;

Fig. 6B shows exemplary 9-N alkylation compounds in accordance with the invention;

Fig. 7A shows a synthetic scheme for introduction of functional groups at the X3 position for interaction with the small hydrophobic binding site;

Fig. 7B shows a second synthetic scheme for introduction of functional groups at the X3 position for interaction with the small hydrophobic binding site;

Figs. 8A and B show synthetic schemes for introduction of variations at the X2 site;

Fig. 9 shows a synthetic scheme for production of binding moieties in accordance with the invention;

Fig. 10 shows a synthetic scheme for introducing variations in the length and character of the bridge between the purine and the phenyl groups in PU family compounds;

Fig. 11 shows dimerization or the addition of targeting moieties;

Fig. 12 shows a schematic methodology for assessing comparative affinities of binding moieties;

Fig. 13 shows the specific structures of PU1-PU4;

Figs. 14A-C show the antiproliferative effect of PU3 on breast cancer cell lines;

Fig. 15 shows X₁ substituents of compounds tested in one example;

Figs. 16A and B show a procedure for modifying PU3 and using biotinylation for immobilization;

Figs. 17A and B show degradation of Akt protein by PU3 and PU24FCI;

Figs. 18 A and B show the correlation between growth arrest, Her2 total protein degradation and Hsp90 binding efficacy; and

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Fig. 19 shows Her2 degradation with PU24FC1.

Detailed Description of the Invention

The present invention relates to small molecules which bind preferentially and with an affinity greater than ADP and less than geldanamycin to the N-terminal pocket of one or more members of the HSP90 family of proteins, for example, to one or more of hsp90 α or β , Grp94 and Trap-1, or proteins with similar pockets, for example DNA gyrase and MutL. As used in the specification and claims of this application, the term "N-terminal pocket of HSP90" refers to the pocket to which geldanamycin and other ansamycin antibiotics bind, and which is occupied by ATP/ADP.

Fig. 1 shows aligned sequences comparing the structures of each of the known members of the HSP90 family of proteins. In the case of hsp90 α , the five potential binding sites are:

- (a) a top binding site comprising Lys112;
- (b) a second top binding site comprising Lys58, Asp54, Phe138 backbone, Gly 135 and Asn106;
- (c) a bottom binding site comprising Asp93, Ser52 and Asn51;
- (d) a hydrophobic lateral binding site comprising Val150, Met98, Val186, Phe138, Leu107, Leu103, Val186 and Trp162; and
- (e) a small hydrophobic bottom binding site comprising Thr184, Val186, Val150 and Ile91.

As indicated in Fig. 1, the N-terminal pocket, while highly conserved in HSP90 family of proteins, does have differences between the various members. The binding moieties of the invention exploit these differences to provide compositions which afford differential degradation of kinases and other signaling proteins. As used in the specification and claims of this application, the term "differential degradation" refers to either degradation of one kinase or signaling protein to a greater extent than another kinase or signaling protein, where both would be degraded in the presence of geldanamycin, or to the degradation of a kinase or signaling protein to a different product than would be obtained in the presence of geldanamycin.

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The size and shape of the N-terminal binding pocket are described in Stebbins et al., "Crystal Structure of an Hsp90-Geldanamycin Complex: Targeting of a Protein Chaperone by an Antitumor Agent" *Cell* 89: 239-250 (1997). There it is observed that "the pocket can be described as a flat-bottomed cone; it is about 15 Å deep, 12 Å in diameter near its surface, 8 Å midway down, and wide enough at the bottom to hold three water molecules. The binding moieties in the pharmaceutical compositions of the invention are designed to fit within this pocket, and to interact with a plurality of the potential binding sites within the N-terminal pocket.

Fig. 2 shows a three dimensional drawing of the pocket with a binding moiety of the type shown in Fig. 4 disposed therein to assist in visualizing the molecular design process. Figs. 3A and B show the conformation of a designed binding moiety based on a purine nucleus (PU3) inside the hsp90 pocket, as determined by x-ray crystallography, inside the pocket, and without the pocket shown, respectively. The opening 10 of the pocket is disposed near the bottom of the figure. The pocket itself has four arms, arm 11 which is nearest the opening 10 and which include the bottom binding site. As reflected in the Fig. 4, the interior of this arm interacts with the amino substituent of the purine nucleus.

Second arm 13 connects to the first arm at an angle of about 95-110°C. Third arm 12 connects to the first arm 11 at an angle of about 100-120°. These two arms contain the two binding sites and are generally hydrophilic in nature. The substituent X₂ (Fig. 4) fits within one of these two arms, resulting in the bending of the binding moiety into the characteristic C-shape. The ability to adopt this C-shape must be taken into account when selecting molecules to act as binding moieties. As noted in Fig. 2, the separation between arms 11 and 13 (9.5 to 11 Å) is greater than the separation between arms 11 and 12 (8-9 Å). Thus, selection of a longer substituent (X₂) may favor insertion of the molecule into arm 13 over arm 12. The fourth arm 14 contains the lateral hydrophobic pocket, and receives the substituent X₃ (Fig. 4). This pocket has a volume of $\approx 100 \text{ Å}^3$ and is generally hydrophobic in character. Thus, the substituent X₁ (Fig. 4) is selected to fill this pocket and to interact with the hydrophobic residues.

In a first embodiment of the invention, the binding moiety is a complete molecule. Fig. 4 shows an exemplary structure of this type based on a purine nucleus,

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and indicates the interactions between the substituents and the binding sites within the N-terminal binding pocket. In Fig. 4, the substituent X_1 may be any hydrophobic chain (linear, branched aliphatic, aromatic, acyclic or cyclic, containing C, H, N, O and/or S atoms that fits within the pocket); X_2 is from 1 to five hydrogen acceptor functionalities such as OR, OCOR, NCOR and the like that fits within the pocket and optionally plus an electron donating group able to enhance the activity of such groups, such as halogen, and X_3 is any small size substituent to fit within the pocket, for example a group such as fluorine. In the embodiment of the invention, a structural difference between the molecules of the invention and geldanamycin antibiotics and derivatives is the interaction between these binding moieties and the hydrophobic lateral binding site and small hydrophobic binding site. As observed in Stebbins et al., these sites are located at the bottom of the pocket, and geldanamycin does not fill these portions. Rather, the methyl group of geldanamycin extends only partly into the lateral pocket, leaving room at the bottom of the pocket for water molecules. In contrast, in compositions of the invention, the substituents may be selected to fill at least the hydrophobic lateral binding site in the pocket. As used in this application, the term "fill" refers to the occupancy of a binding site to an extent such that there is not room remaining to accommodate a water molecule.

Additional examples of binding moieties within the scope of the invention that are designed to fit within the N-terminal pocket of HSP90 proteins and interact with the pockets with affinity between that of ADP and geldanamycin for at least one member of the HSP90 family are shown in Fig. 5. In these compositions, X_1 is a hydrophobic chain (linear, branched aliphatic, aromatic, acyclic or cyclic, containing C, H, N, O and/or S atoms that first within the pocket), or COR, where R is a hydrophobic chain; X_2 is from 1 to 5 hydrogen acceptor/donor functionalities, which may be the same or different; X_3 is a small size group such as alkyl (saturated, unsaturated, cyclic or linear), alkoxy (for example methoxy or ethoxy), halogen or SR (where R is methyl or ethyl); X_4 is H or as X_3 , X_5 is H or as X_1 ; X_6 is $-NH_2$, $-OR$ (R being H or alkyl), or $-CONH_2$ or a similar hydrogen donor functionality and R_1 is H or alkyl. These molecules are all capable for folding into a C-shaped configuration and interact with at least three of the potential binding sites, including the hydrophobic lateral site. It will be appreciated that other compounds with different nuclei but comparable overall volume (~ 200 - 500 \AA^3) and

dimensions can be designed in the same manner and fall within the scope of the present invention.

The compositions having structures indicated as the PU family in Fig. 5 can be synthesized using the procedure outlined in Fig. 6A. A carboxylic acid starting material is converted to an acid fluoride by reaction with cyanuric fluoride in pyridine/DCM or to an acid chloride by reacting with SOCl_2 . This acid halide is reacted with an amino-substituted pyrimidine (such as 4,5,6-triaminopyrimidine sulfate in DIEA/DMF or 2,4,5,6-tetraaminopyrimidine sulfate in aqueous NaOH, or either sulfate with K_2CO_3 in DMF) to produce an intermediate product (PY-A, PY-B, PY-C in Fig. 1) which undergoes ring closure to produce substituted 8-benzyl purine derivatives (PU-A, PU-B, PU-C) which are useful compositions in accordance with the invention. If desired, a alkylation reaction, such as a Mitsunobu alkylation, may be performed to add an alkyl group, R, (with or without functional substituents) to the nitrogen at the 9-position. Exemplary compounds made by 9-N alkylation of the PU family precursor PU-C are shown in Fig. 6B.

Figs. 7 A and B show synthetic schemes for introduction of functional groups at the X3 position for interaction with the small hydrophobic binding site. Figs. 8A and B show synthetic schemes for introduction of variations at the X2 site.

Fig. 9 shows a synthetic scheme for synthesis of the RD class of compounds from Fig. 5. Other classes shown in Fig. 5 can be made using comparable synthetic approaches.

The compositions of the invention may be used directly to provide therapeutic benefits in the treatment of cancers and other diseases. As illustrated in the examples below, the compositions of the invention have been shown to induce degradation in Her2 kinase in SKBr3, MCF-7 and BT474 breast cancer cells. In addition, compositions of the invention have been shown to be effective at causing Rb-positive SKBr3 and MCF-7 breast cancer cell lines to undergo G1 arrest, and to have antiproliferative effects against these cell lines and BT474, MDA-MB468 and prostate cancer cell lines TSUPr and LNCaP.

The compounds of the PU family were tested for binding to Hsp90, degradation of Her2 total protein and for their antiproliferative effect. Table 1

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summarizes the influence of the 9-N chain on this activity. Compounds of the PU family (Fig. 6) not listed in the table were either inactive or insoluble.

Compounds of the PU family were modified by addition of a fluorine group as substituent X3 (C-2 fluorination) using the reaction protocol of Fig. 7B. The results are summarized in Table 2. As shown, introduction of a fluorine substituent increased the activity of compositions with common X1 groups, and enhanced the activity/solubility of compositions with X1 substituents for which activity was not reported in Table 1.

Variations in activity were also observed when additional halogen substituents were added to the phenyl ring using the reaction scheme shown in Fig. 8B. As shown in Table 3, introduction of a halide at one, but not both of the positions not occupied by methoxy groups in PU3 led to an increase in activity relative to PU3.

Compounds were prepared with the chlorine substituent as in compound PU3PhCl and the fluorine substituent as in PU24F or PU29F. As shown in Table 4, these compounds were the most active of those tested so far.

Another location of potential variation in the PU family of molecules is in the length and character of the bridge between the purine and the phenyl groups. Fig. 10 shows a synthetic scheme for achieving such variation. Compound 69 was tested and was found to have weak binding to hsp 90-beta, but no detectable binding to hsp 90-alpha. This is the reverse of the binding selectivity observed for Compound 18. This provides for selectivity between these two proteins.

As an alternative to the use of the compositions of the invention individually, coupled-compositions in which the hsp-binding nucleus as shown in general structures in Fig. 5 are derivatized by coupling to targeting moiety selected to specifically bind to a protein, receptor or marker found on a target population of cells or it may be dimerized. (See Fig. 11) The targeting moiety may be a hormone, hormone analog, protein receptor- or marker-specific antibody or any other ligand that specifically binds to a target of interest. Particular targeting moieties bind to steroid receptors, including estrogen and androgen and progesterone receptors, and transmembrane tyrosine kinases, src-related tyrosine kinases, raf kinases and PI-3 kinases. Specific tyrosine kinases include HER-2 receptors and other members of the epidermal growth factor (EGF)

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receptor family, and insulin and insulin-like growth factor receptors. Examples of specific targeting moieties include estrogen, estradiol, progestin, testosterone, tamoxifen and wortmannin. Targeting moieties may also be antibodies which bind specifically to receptors, for example antibodies which bind to Her2 receptors as disclosed in International Patent Publications Nos. WO96/32480, WO96/40789 and WO97/04801, which are incorporated herein by reference.

Because of their ability to bring about the degradation of proteins which are essential to cellular function, and hence to retard growth and/or promote cell death, the hsp-binding compounds of the invention, with or without a targeting moiety, can be used in the therapeutic treatment of a variety of disease conditions. A suitable therapeutic is one which degrades a kinase or protein that is found in enhanced amounts or is mutated in disease-associated cells, or on which the viability of such cells depends. The general role of HSP90 proteins in maintaining malignancy in most cancer cells points to the importance of this target in the development of anticancer agents. Thus, the therapeutic small molecules of the invention provide a novel modality for the treatment of all cancers that require or are facilitated by an HSP90 protein. For example, the compositions of the invention can be used in the treatment of a variety of forms of cancer, particularly those that overexpress Her2 or mutated or wild type steroid receptors, or that lack functional RB protein. Such cancers may include but are not limited to breast cancer and prostate cancer. In addition, the compositions of the invention can be used in the treatment of other diseases by targeting proteins associated with pathogenesis for selective degradation. Examples of such targetable proteins include antigens associated with autoimmune diseases and pathogenic proteins associated with Alzheimer's disease.

The compositions of the invention are suitably utilized to degrade specific proteins which are associated with the disease state or condition of concern. Because the compositions of the invention can be selected to degrade specific kinases or signaling proteins, a suitable therapeutic is one which degrades a kinase or protein that is found in enhanced amounts in diseased cells. Thus, for example, the selection of a binding moiety which degrades Her2 kinase, but not other HER kinases or Raf kinase would be suitable for treatment of Her2- positive breast cancer. The examples below provide guidance on the selection of specific compounds based on the specificity observed *in vitro*. In

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addition, screening techniques are described below to facilitate the evaluation of structures for binding affinity and differential degradation.

The compositions of the invention are administered to subjects, including human patients, in need of treatment, in an amount effective to bring about the desired therapeutic result. A suitable administration route is intravenous administration, which is now commonly employed in chemotherapy. In addition, because the compositions of the inventions are small soluble molecules, they are suitable for oral administration. The ability to use an oral route of administration is particularly desirable where it may be necessary to provide treatment of a frequent, for example a daily schedule. The amount of any given composition to be administered, and the appropriate schedule for administration are determined using toxicity tests and clinical trials of standard design, and will represent the conclusion drawn from a risk benefit analysis. Such analyses are routinely performed by persons skilled in the art, and do not involve undue experimentation.

As an alternative or adjunct to the design of molecules for use as binding moieties in accordance with the invention, we have developed a fast screening assay which can be used to test compounds for their binding affinity to the N-terminal pocket of HSP90, or to test proteins for the presence of an HSP90 type of pocket. As summarized in Fig. 12, the surface of a solid substrate (such as the bottoms of 96 well microtiter plates) are functionalized with N-oxysuccinimydyl functionalities. These react in isopropanol with amino-functionalized GM (or other ansamycin antibiotic). The amount of GM bound to the wells can be assessed by spectrophotometric absorbance measurements at 345 nm. When the bound GM is incubated with rabbit reticulocyte lysates, enough hsp90 is captured to be detectable by colorimetric methods, although more specific detection methods are preferred. For example, bound hsp90 can be detected using a labeled anti-HRP antibody. To assay for HSP90 binding efficacy, a test compound is added to the wells with the rabbit reticulocyte lysate (or other hsp90 source) and any differences in the amount of captured hsp are noted. If the test compound binds hsp90 with greater affinity than GM, it will compete with the immobilized GM and result in a reduction in the amount of hsp90 captured. By varying the member of the HSP90 family used in the assay, these same plates can be used to evaluate differences in

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specificity of test compounds. The plates can also be used to screen protein/peptide libraries for proteins which possess an HSP90 type of binding pocket.

A test was also designed which allows identification of compounds which interact differentially with Hsp90-alpha and Hsp90-beta. In this test, geldanamycin (or other strong non-discriminating hsp90 binder such as other ansamycin antibiotics or radicicol) is modified as necessary and affixed to a solid support, for example beads. An Hsp90 protein preparation containing both the alpha and the beta form is incubated with the support in the presence or absence of a compound to be evaluated. If the compound binds to the Hsp90 protein, it competitively inhibits the binding of the protein to the solid support. After washing, the material bound to the support is eluted and the eluate is separated on an SDS/PAGE gel and visualized by immunoblotting with anti-Hsp90-alpha and anti-Hsp90-beta to determine the amount of material bound. Alternatively, if quantitative amounts of the Hsp90 protein preparation or known ratios of alpha to beta forms of Hsp90 are used in the first instance, the unbound material can be analyzed by a similar technique.

Candidate compounds in accordance with the invention can also be evaluated for their ability to deplete or induce proteins which are found in enhanced amounts in cancer cells or on which the cells depend for viability (for example Her2, Her3, Raf-1, ER, Rb, cdk-4, Hsp90, Trap-1, and Grp94) in a panel of cancer cell lines using the cell-based assay of Stockwell et al., *Chem Biol.* 6: 71-83 (1999). In this assay, cells are grown at the bottom of a well and fixed. The fixed cells are probed for the presence of a specific primary antigen (the oncogenic protein) using a specific primary antibody in solution. A secondary antibody covalently linked to horseradish peroxidase is added, and the resulting complex is detected through the chemiluminescent reaction caused by the addition of luminol, hydrogen peroxide and an enhancer such as p-iodophenol. Differences in the detected luminescence levels in the presence of a drug indicate activity of that drug with respect to the degradation or induction of the targeted protein.

Drugs may also be assayed based upon observed very characteristic changes in cell morphology. MCF-7 cells exposed to PU3 flatten, increase in size and have distinct cellular boundaries. The increase in size is mostly due to an abundance in

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cytoplasm. These morphological changes are characteristic of mature epithelial differentiation and reversal of transformation.

As a third alternative, Immunofluorescence (IF) and Hematoxylin and Eosin stain (H&E) may be used to assess drug effectiveness. Cells were plated on 8 well chamber slides, Fisher Scientific) and seeded for 24hrs. Drugs or vehicle were added for 5 days after which, for IF the slides were washed twice with ice-cold PBS and fixed with methanol and acetone solution (1:1) for 15 sec. Fixed monolayers were washed with distilled water and blocked with 5% BSA in PBS solution. After blocking, cells were incubated with the primary antibody (anti-MFMG, Chemicon, 1:100 in 5% BSA in PBS) at 37°C and washed 3 times with 1% BSA in PBS, followed by incubation with a rhodamine-labeled secondary antibody for 1hr at 37°C. Nuclei were stained with DAPI at 1mg/ml. For H&E, the cells were fixed with paraformaldehyde (4%) for 10min at RT and stained according to standard H&E staining protocols. The induction of G1 arrest by a variety of manipulations is sufficient for induction of expression of some milk proteins. However, only ansamycins, the HDAC inhibitor SAHA and the hsp90 binding molecules cause extensive morphological and biological changes.

For use in the therapeutic method of the invention, the compositions of the invention are formulated in a pharmaceutically acceptable carrier. For injectable formulations, this may be a sterile solution (for example sterile saline), or the compounds may be formulated in a lipidic carrier. For oral formulations, the compositions of the invention may be packaged in convenient dosage unit form, for example as tablets or capsules with suitable excipients, or as a liquid formulation. In the latter case, the liquid pharmaceutical will suitably include a flavorant to enhance palatability, and may also include colorants and other conventional additives.

EXAMPLE 1

Compositions having general formula 1 were synthesized according to the scheme shown in Fig. 6A. To generate acid fluorides, to a solution of the phenyl acetic acid derivative (3 mmol) in 15 mL of CH₂Cl₂ (under inert atmosphere) were added 1.5 equivalents of cyanuric fluoride and pyridine (3 mmol). The mix was stirred for 1.5h at room temperature, after which, 30 more mL of CH₂Cl₂ were added. The resulting solution

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was washed once with 0.5 mL water, and the crude acid fluoride material resulted from the removal of the solvent was used in the next step.

The acid fluoride was used for the generation of PY-A, PY-B and PY-C derivatives as follows: To 675 mg (2.8 mmol) of 4,5,6-triaminopyrimidine sulphate in 1.5 mL of DMF were added 1.5 mL (9 mmol) of DIEA, the acid fluoride (obtained as described above) dissolved in 5 mL CH_2Cl_2 and a catalytic amount of DMAP (0.28 mmol). The reaction mixture was stirred under argon for 1 hr. The solid that formed was filtered off and the filtrate was concentrate to dryness. To the residue material was added 10 mL EtOAc and the precipitate formed was washed several times with EtOAc and CH_2Cl_2 to give a slightly yellow material in 60-80% yield. This material was used for the next step. If higher purity is desired, flash chromatography using EtOAc: MeOH=7:1 (1% TEA) can be performed.

From the PY compounds, PU-A, PU-B and PU-C derivatives were generated as follows: To 500 mg (1.5 mmol) of 5-acylated-4,5,6-pyrimidine in 13 mL MeOH was added 13 mL solution of 25% NaOMe in MeOH and the resulting solution was refluxed for 5hrs. After cooling, 5mL of water was added and the solution was concentrated to 5 mL. The resulting aqueous solution was extracted with 4x10 mL THF and 2x10 mL EtOAc. The combined organic layers were dried with MgSO_4 , concentrated to dryness to give a slightly yellow solid in 80-90% yield. If higher purity is desired the resulting product can be added to a silica gel column and eluted with CH_2Cl_2 :EtOAc:MeOH=4:4:1.

Substituents were added to the 9-N position using a Mitsunobu alkylation as follows: To 0.16 mmol of purine derivative in 5mL toluene and 1 mL CH_2Cl_2 were added 2.2 equivalents PPh_3 and 1.3 equivalents of ROH followed by 5 equivalents of DEAD. The reaction was monitored by TLC and proceeded in 15 min to 1 hr. The mixture was applied to a ISCO CombiFlash system (silica gel column) and eluted with a gradient of CH_2Cl_2 /acetone to give 30-75% isolated product. This procedure was carried out using the 40 unbranched and branched, linear and cyclic, saturated and unsaturated primary and secondary alcohols shown in Fig. 6. In these syntheses, only two alcohols, neopentyl alcohol and cyclohexanol, were found to give no product in this reaction.

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Four of the compounds (PU-A-4, PU-B-4, PU-C-4 and PU-C-15) were used in initial tests. For convenience, these four structures are referred to as PU1, PU2, PU3 and PU4, respectively as shown in Fig. 13.

EXAMPLE 2

The human cancer cell lines MCF-7, SKBr3 and MDA-MB-468 were obtained from the American Type Culture Collection (Manassas, VA) and maintained in 1:1 mixture of DME:F12 supplemented with 2 mM glutamine, 50 units/mL penicillin, 50 units/mL streptomycin and 5% (for MCF-7 and MDA-MB-468) or 10% (for SKBr3) heat inactivated fetal bovine serum (Gemini Bioproducts) and incubated at 37°C in 5% CO₂.

For assays on protein levels, cells were grown to 60-70% confluence exposed to drugs or DMSO vehicle for the indicated time periods. Lysates were prepared using 50mM Tris pH=7.4, 2% SDS and 10% glycerol lysis buffer. Protein concentration was determined using the BCA kit (Pierce Chemical Co.), according to the manufacturers instructions. Clarified protein lysates (20-50mg) were electrophoretically resolved on denaturing SDS-PAGE, transferred to nitrocellulose and probed with the following primary antibodies: anti-Her2 (C-18), -Her3 (C-17), -Raf-1, -cyclin D1, -Rb (C-15) (Santa Cruz), anti-hsp90, -hsp70, -ER (Stressgen), anti-Trap-1 (MSK81), anti-b-actin, -tubulin (Sigma), anti-PI3K (p85) (Upstate Biotechnologies).

To determine antiproliferative indices, growth assays were performed by seeding 10,000 cells per well in 6-well dishes and incubating for 24hrs before drug treatment. Drugs or vehicle were administered as outlined for each experiment, and cells were incubated for the time periods depicted and then counted by coulter counter.

Cell cycle distribution was assayed according to Nusse et al with a Becton Dickinson fluorescence activated cell sorter and analyzed by Cell Cycle Multi-cycle system (Phoenix Flow System, San Diego, CA).

Experiment 1:

MCF-7 cells were treated with varying concentrations (0, 10, 50, 100 or 250 µM) of PU3 or the control purine Ad-But for 24 hours and then lysed. Levels of Hsp90, Trap1 and Hsp70 were evaluated by Western blotting. The results showed that

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PU3, like GM, increases the cellular levels of Hsp90 and Hsp70. Treatment of cells with PU3 also induces a protein band that migrates more rapidly than Trap-1 and that is recognized by an anti-Trap-1 antibody. Although the identity of this protein band is unknown, its appearance seems to be a marker of cellular exposure to Hsp90 inhibitors. Ad-But had no effect on the studied protein levels at identical concentrations.

Experiment 2:

Cell cultures of MCF-7, SKBr3 and MDA-MB-468 cells were treated with PU3 at one of several concentrations or a DMSO control to test for the occurrence of antiproliferative effects. Figs. 14A-C show the results of cell number as a function of time. As shown, PU3 clearly inhibits growth of these breast cancer cell lines.

Experiment 3:

Cell cultures of SKBr3 cells were treated with PU1, PU2, PU3 or PU4 at concentrations of 0, 10, 50, 100 and 250 μ M. After 24 hours, the cells were lysed and the amount of Her2 and Raf-1 in the cells was evaluated. Although the compounds are structurally very similar, they were found to have different efficacy at promoting degradation of the two proteins. PU1 showed little degradation of either protein at any of the concentrations tested, while PU2 degraded Her2 but not Raf-1 at 250 μ M. PU3 degraded Her2 at concentrations over 100 μ M but resulted in only partial loss of Raf-1 at 250 μ M. PU4 degraded Her2 at concentrations greater than 50 μ M and Raf-1 at a concentration of 250 μ M.

Experiment 4:

To test the time course of degradation, MCF-7 cells were treated with 100 μ M PU3. Aliquots of cells were recovered at 1.5, 3, 6, 12 and 24 hours, and the amount of Her2, Raf-1, Hsp90 and Trap-1 was evaluated. The compound was shown to induce rapid degradation of Her2 (with more than 50% being lost within 3 hours of administration) and significant synthesis of Hsp90 and Hsp70 within the 24 hour period. The amount of Raf-1 did not change significantly during this time.

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Experiment 5:

Cell cultures of MCF-7 cells were treated with PU1, PU2, PU3 or PU4 at concentrations of 0, 10, 50, 100 and 250 μ M. After 24 hours, the cells were lysed and the amount of Her2 in the cells was evaluated. Although the compounds are structurally very similar, they were found to have different efficacy at promoting degradation of Her2. PU1 showed only partial degradation at concentrations in excess of 50 μ M, while PU2 degraded Her2 completely at 250 μ M. PU3 substantially degraded Her2 at concentrations over 10 μ M. PU4 substantially degraded Her2 at concentrations greater than 100 μ M.

Experiment 6:

Cell cultures of MCF-7 cells were treated with PU1, PU2, PU3 or PU4 at concentrations of 0, 10, 50, 100 and 250 μ M. After 24 hours, the cells were lysed and the amount of estrogen receptor in the cells was evaluated. Although the compounds are structurally very similar, they were found to have different efficacy at promoting degradation of estrogen receptors. PU1 showed essentially no degradation at the concentrations tested, while PU2 degraded estrogen receptor completely at 250 μ M. PU3 partially degraded estrogen receptor at concentrations over 50 μ M. PU4 substantially degraded Her2 at concentrations greater than 100 μ M.

Experiment 7:

Cultures of SKBr3 and MCF-7 cells were treated with varying concentrations (0, 10, 50 100, or 250 μ M) of PU3 for 24 hours. The cells were lysed and levels of Her2, Raf1, Her3, estrogen receptor (ER), p85 (PI3K), tubulin and actin were analyzed by western blotting. Reduction in the amount of those signaling proteins which depend on hsp90 for their function (Her2, Raf1, Her3 and ER) was observed. No effect was observed on PI3K, tubulin or actin, proteins which are involved in other signaling pathways.

EXAMPLE 3

The experiments of Example 2 demonstrate the ability of synthetic compounds PU1, PU2, PU3 and PU4 in accordance with the invention to provide

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differential degradation of kinases HER3, estrogen receptor (ER), Her2, Raf, Rb and p85 (as reference protein which should not be, and is not degraded). Several important observations can be made from this data.

PU3 destabilizes the estrogen receptor complex and induces dose-dependent degradation of the protein. PU3 also causes a rapid decrease in Her2 levels and causes accumulation of a lower-molecular weight (170 kDa versus 180 kDa mature protein) HER-2 band, also seen for GM, that is thought to be incompletely glycosylated Her2 which is sequestered *in vivo* in the endoplasmic reticulum. Levels of Raf and Her3 are less sensitive to PU3, but are degraded at higher concentrations. PU2 also induces a lower molecular weight Rb band which is thought to be hyperphosphorylated Rb.

In contrast, PU2 showed substantial degradation of Her2 without accumulation of the lower molecular weight band and was less effective for degradation of Raf kinase. The lower molecular weight band is also absent when PU1 is used as the binding moiety. PU2 is less effective at degrading ER, while PU1 has had substantially no effect on ER. These differences provide evidence suggesting that PU3 binds to a different member of the HSP90 family from PU1 and PU2, thus accounting for the different specificity. PU4 also shows greater ability to degrade Her2 than ER or Raf.

EXAMPLE 4

Six compounds of the PU-C family (X4= H, X2=1, 2, 3 OMe, X3=H and with variable X1 groups as indicated in Fig. 15) were prepared and tested for their effect on Her2, HER3 and TRAP-1 in MCF-7 cells. The cells were treated for 24 hours with 10, 50, 100 and 250 μ M of each compound. The nature and size of the X1 group has a significant effect on the activity of the compound. Large, hydrophobic groups which will better fill the N-terminal pocket show the ability to degrade Her2, but not Her3. In addition, for these compounds there is a correlation between the ability to degrade Her2 and induction of Trap-1 synthesis.

On the other hand, the compounds with big, rigid and somewhat polar X1 groups interact less favorably, and do not significantly degrade Her2 at the concentrations tested.

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EXAMPLE 5

PU3 was linked to Biotin through the middle OH in accordance with the synthetic scheme shown in Figs. 16A and B. PU3-biotin was immobilized on Sepharose-Streptavidin beads. These beads were used to show the interaction of PU3 with the Trap-1 and Hsp90alpha.

EXAMPLE 6

PU3 was tested for its safety and potency in MCF-7 xenografts. The compound was administered IP up to 500 mg/kg without showing significant signs of toxicity. Mice with established tumors were treated with a single dose of PU3 at doses of 50, 100 and 500 mg/kg i.p. Control mice were treated with DMSO alone. Mice were sacrificed 12 hours post-treatment. For immunoblotting, tumor tissue was homogenized in 2% SDS lysis buffer (pH 7.4). In all cases a reduction in Her2 levels was observed in the immunoblotting results. No change was observed in levels of PI3 kinase (p85).

EXAMPLE 7

Several 9-Alkyl-8-benzyl-9H-purin-6-ylamines (4-46) were created by 9-N-alkylation of PU-C (Fig. 6B). Additionally, the 2-amino group of 3 (Fig. 7B) was converted to fluorine via a diazotization reaction in non-aqueous media using *tert*-butyl nitrite (TBN) in HF/pyridine. The resulting purine was alkylated to give the 2-fluoro-9-alkyl-8-(3,4,5-trimethoxy-benzyl)-9H-purin-6-ylamines 47-59. The alkylation was performed using the Mitsunobu reaction, methodology that can accommodate a large array of primary and secondary alcohols. The 2-fluoro derivatives were converted to the 2-alcoxy purines 60 and 61 by refluxing in the corresponding alcohol and NaOMe. The majority of modifications at position 2 of the purine moiety commenced with (3). Iodine was introduced in that position using isoamyl nitrite in diiodomethane to give 62. The 2-iodo derivative was transformed to the cyano-derivative 63 with tri-*n*-butylcyanostannane and tetrakis(triphenylphosphine)palladium(0) in DMF and to the vinyl-derivative 64 with vinyltributyltin and (MeCN)₂PdCl₂.

The benzyl moiety was enriched in electron-donating groups in order to increase its interaction with the pocket lysine. Chlorine and bromine were added via a

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radicalic reaction using *t*-butyl hydroperoxide and the corresponding acid halide. In the case of chlorine only monosubstitution was observed (65), while bromine gave the monosubstituted (66) and a small amount of di-bromosubstituted product (67). (Fig. 8B)

The nature and length of the bridge between the purine and the phenyl ring were additionally modified. As starting material we utilized the 8-Br-adenine. This was reacted at high temperature with the aniline-, phenol- or benzyl-derivative to give the corresponding products 68-70. (Fig. 10)

The assembly of the fully substituted derivatives 71 and 72 commenced with 3. Chlorine was added initially via the radicalic reaction described above to give 73. Subsequently, the 2-amino group was transformed to fluorine and the resulting purine (74) was alkylated via the Mitsunobu reaction.

Specific reactions for various compounds are set forth below:

3 (DAAC) :8-(3,4,5-Trimethoxy-benzyl)-9H-purine-2,6-diamine: To trimethoxyphenyl acetic acid (1.0 g, 4.4 mmol) in DCM (20 mL) (under inert atmosphere) was added cyanuric fluoride (371 mL, 4.4 mmol) and pyridine (356 mL, 4.4 mmol). The mixture was stirred for 1 h at room temperature, after which, an additional 30 mL of DCM was added. The resulting solution was washed once with water (5 mL), and the acid fluoride resulted from the removal of the solvent was taken up in DMF (10 mL) and used in the next step. Separately, 2,4,5,6-tetraaminopyrimidine sulfate (0.9 g, 3.8 mmol) was dissolved in water (40 mL) that contained NaOH (456 mg, 11.4 mmol). The resulting solution was heated to 70°C and the acid fluoride was added dropwise over a 20 minute period. The reaction mixture was stirred for 1.5 h at 70°C and then concentrated to dryness. To the crude material was added MeOH (16 mL) and a 25% solution of NaOMe in MeOH (16 mL) and the resulting solution was heated at 90°C for 18 h. Following cooling, the pH of the solution was adjusted to 7 by addition of concentrated HCl. The aqueous solution was removed and the crude taken up in DCM (100 mL) and MeOH (50 mL). The undissolved solids were filtered off and the product (470 mg, 37%) was purified on a silica gel column with EtOAc:DCM:MeOH at 4:4:2.

FAC: 2-Fluoro-8-(3,4,5-trimethoxy-benzyl)-9H-purin-6-ylamine: To 3 (500 mg, 1.5 mmol) was slowly added a 70% solution of HF in pyridine (2 mL), pyridine (8 mL),

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followed by *t*-butyl nitrite (200 mL, 2.0 mmol). The mixture was stirred for 2 h and then quenched overnight with 8 g CaCO₃ in water (15 mL) and MeOH (10 mL). The solution was concentrated to dryness and the resulting crude was taken up in MeOH (30 mL) and DCM (10 mL). The insoluble solids were filtered off and the solvent was removed to give the crude product. This was purified on a silica gel column eluting with hexane:DCM:EtOAc:MeOH at 10:5:5:0.75 (240 mg, 50%).

The alkylation was performed via a Mitsunobu reaction as described before (ref). Essentially, to FAC in toluene:DCM at 5:1 was added 1.3 eq. of the corresponding alcohol, 2 eq. PPh₃, 3 eq. di-*tert*-butyl azodicarboxylate and the resulting solution was stirred at room temperature for 10min to 1 h (conversion was monitored by TLC) to give:

48 (PU3F): 2-Fluoro-9-butyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 60% yield.

52 (PU29F): 2-Fluoro-9-(2-isopropoxy-ethyl)-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 86% yield.

54 (PU47F): 2-Fluoro-9-pent-4-enyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 86% yield.

59 (PU20F): 2-Fluoro-9-(tetrahydrofuran-2-ylmethyl)-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 53% yield.

50 (PU44F): 2-Fluoro-9-(2-methoxy-ethyl)-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 28% yield.

47 (PU43F): 2-Fluoro-9-propyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 66% yield.

57 (PU24F): 2-Fluoro-9-pent-4-ynyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 76% yield.

53 (PU8F): 2-Fluoro-9-but-3-enyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 42% yield.

55 (PU48F): 2-Fluoro-9-but-3-ynyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 36% yield.

56 (PU49F): 2-Fluoro-9-pent-3-ynyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 25% yield.

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58 (PU16F): 2-Fluoro-9-cyclobutylmethyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 33% yield.

51 (PU26F): 2-Fluoro-9-[(*S*)-2-methyl-butyl]-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 68% yield.

49 (PU21F): 2-Fluoro-9-pentyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: 65% yield.

PU24DA: 9-Pent-4-ynyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purine-2,6-diamine: To DAAC (200 mg, 0.61 mmol) of in toluene (20 mL) and DCM (4 mL) was added PPh₃ (330 mg, 1.3 mmol), 4-pentyne-1-ol (75 mL, 0.8 mmol) and di-*t*-butyl azodicarboxylate (600 mg, 2.5 mmol). The mixture was stirred at room temperature for 2 h. The product was isolated by column chromatography on a silica gel column eluting with hexane:CHCl₃:EtOAc:EtOH at 10:8:4:4 (120 mg solid, 49%).

62 (PU24I): 2-Iodo-9-pent-4-ynyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: To PU24DA (50 mg, 0.13 mmol) was added CH₂I₂ (2.5 mL) and isoamyl nitrite (100 mL, 0.78 mmol) and the resulting solution was heated at 80°C for 1 h. After cooling, the solution was concentrated and then added to a silica gel column. The product was isolated (40 mg, 61%) using eluent hexane:CHCl₃:EtOAc:EtOH at 10:4:4:0.75.

63 (PU24CN): 2-Cyano-9-pent-4-ynyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: A solution of PU24I (20 mg, 0.04 mmol), Pd(PPh₃)₄ (7 mg, 6.3x10⁻³ mmol) and tributyltin cyanide (14 mg, 0.043 mmol) in dry DMF (2.5 mL) was heated at 180°C for 6 h. Following cooling and removal of the solvent, the product (13 mg, 81%) was isolated by column chromatography (hexane:CHCl₃:EtOAc:EtOH at 10:8:4:0.75).

64 (PU24V): 9-Pent-4-ynyl-8-(3,4,5-trimethyl-benzyl)-2-vinyl-9*H*-purin-6-ylamine: A solution of PU24I (20 mg, 0.04 mmol), (MeCN)₂PdCl₂ (2x10⁻³ mmol) and tributylvinyltin (12.5 mL, 0.041 mmol) in dry DMF (3 mL) was heated at 100°C for 1 h. Following cooling and removal of the solvent, the product (15 mg, 92%) was isolated by column chromatography (hexane:CHCl₃:EtOAc:EtOH at 10:8:4:1).

60 (PU3OMe): 2-Methoxy-9-butyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: To a solution of PU3F (20 mg, 0.051 mmol) in MeOH (1 mL) was added a 25% solution of NaOMe in MeOH (1.5 mL). The resulting mixture was heated at 85°C for 2 h. Subsequent to cooling, the mixture was neutralized with 4 N HCl and then concentrated to dryness.

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The crude was purified on a silica gel column with hexane:EtOAc:DCM:MeOH at 10:5:5:1 to give a solid (11 mg, 51%).

61 (PU3OEt): 2-Ethoxy-9-butyl-8-(3,4,5-trimethoxy-benzyl)-9H-purin-6-ylamine: To a solution of PU3F (20 mg, 0.051 mmol) in EtOH (2 mL) was added NaOMe (15 mg, 0.28 mmol) and the mixture was refluxed for 2 h. After cooling and neutralization with HCl, the solution was concentrated to dryness. The product (12 mg, 56%) was purified as described above.

65 (PU3Cl): 9-butyl-8-(2-chloro-3,4,5-trimethoxy-benzyl)-9H-purin-6-ylamine: To PU3 (37 mg, 0.1 mmol) in MeOH (3 mL) was added concentrated HCl (33 mL, 0.4 mmol). The solution was cooled to 0°C and a 90% aqueous solution of *t*-butyl hydroperoxide (44 mL, 0.4 mmol) was added. The resulting solution was refluxed overnight. Following cooling and removal of the solvent, the product (31 mg solid, 72%) was isolated by column chromatography (elute with hexane:EtOAc:DCM:MeOH at 10:5:5:1.5).

66 and 67 (PU3PhBr and PU3PhBr2): To PU3 (37 mg, 0.1 mmol) in MeOH (3 mL) was added a 48% aqueous solution of HBr (45 mL, 0.4 mmol). The mixture was cooled to 0°C, and *t*-butyl hydroperoxide (44 mL, 0.4 mmol) was added dropwise. The resulting solution was stirred for 30 minutes at 0°C and then refluxed for an additional hour. Following cooling and removal of the solvent, the mixture was separated by column chromatography (elute with hexane:EtOAc:DCM:MeOH at 10:5:5:1.5) to give predominantly monobrominated product (31 mg, 69%) and a trace of dibrominated product (2.3 mg, 4.4%). **66 (PU3Br):** 9-butyl-8-(2-bromo-3,4,5-trimethoxy-benzyl)-9H-purin-6-ylamine; **67 (PU3Br2):** 9-butyl-8-(2,6-dibromo-3,4,5-trimethoxy-benzyl)-9H-purin-6-ylamine.

73 (DAACPhCl): To a slurry of DAAC (1 g, 3.0 mmol) in MeOH (50 mL) was added concentrated HCl (1.5 mL, 18 mmol). The resulting solution was cooled to 0°C and a 70% aqueous solution of *t*-butyl hydroperoxide (2.5 mL, 8 mmol) was slowly added. The mixture was stirred for 30 min at 0°C and then refluxed for 20 h. The solvent was removed under high vacuum to give clean product (1.09 g, 98%).

74 (FDAACPhCl): To a 20% solution of HF in pyridine (6.5 mL) was added (under inert atmosphere) pyridine (13.5 mL), followed by DAACPhCl (1 g, 2.7 mmol). The mixture

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was stirred for 5 min and consequently, *t*-butyl nitrite (450 mL, 3.5 mmol) was slowly added. Stirring continued for another 30 min. The reaction was quenched by addition of CaCO₃ (16.5 g) in water (10 mL): MeOH (10 mL) and stirring for 2 h. The solution was concentrated and the resulting slurry was taken up in DCM (50 mL): MeOH (50 mL). The insoluble solids were filtered off and washed with DCM:MeOH at 1:1 (2x25 mL).

Following solvent removal, the product was purified on a silica gel column eluting with hexane:EtOAc:DCM:MeOH at 10:5:5:1 (370 mg solid, 37%).

71 (PU24FCI): 2-Fluoro-9-pent-4-ynyl-8-(2-chloro-3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: (308 mg, 76%).

72 (PU29FCI): 2-Fluoro-9-(2-isopropoxy-ethyl)-8-(2-chloro-3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine:

68 (PAM3): 9-Butyl-N⁸-(3,4,5-trimethoxy-phenyl)-9*H*-purine-6,8-diamine: A mixture of BrAd3 (50 mg, 0.186 mmol) and trimethoxyaniline (120 mg, 0.65 mmol) was heated at 160°C for 30 min. Following cooling, the solid was taken up in DCM (20 mL) and MeOH (5 mL) and any insoluble solids were filtered off. The product (57 mg solid, 83%) was purified on a silica gel column eluting with DCM:EtOAc:MeOH at 7:4:1.

70 (PU3OBn): 9-Butyl-8-(3,4,5-trimethoxy-benzyl)-9*H*-purin-6-ylamine: A solution of benzyl alcohol (250 mL, 2.5 mmol) in 25% NaOMe in MeOH (50 mL) was stirred for 5 min. Following the removal of methanol, BrAd3 (26 mg, 0.096 mmol) and Cu powder (10 mg, 0.16 mmol) was added, and the mixture was heated for 2 min at 180°C. The product was purified on silica gel column with hexane:DCM:EtOAc:MeOH at 10:5:5:1 (5.5 mg, 14%).

69 (PU3OPh): 9-Butyl-8-(3,4,5-trimethoxy-phenoxy)-9*H*-purin-6-ylamine. A mixture of trimethoxyphenol (75 mg, 0.4 mmol), *t*-BuOK (34 mg, 0.3 mmol), Cu powder (10 mg, 0.64 mmol) and BrAd3 (26 mg, 0.1 mmol) was heated for 1 h at 140°C. The product (13 mg solid, 35%) was isolated by column chromatography (hexane:DCM:EtOAc:MeOH at 10:5:5:1).

EXAMPLE 8

The human cancer cell lines MCF-7, SKBr3 and MDA-MB-468 were obtained from the American Type Culture Collection (Manassas, VA) and maintained in

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1:1 mixture of DME:F12 supplemented with 2 mM glutamine, 50 units/mL penicillin, 50 units/mL streptomycin and 5% (for MCF-7 and MDA-MB-468) or 10% (for SKBr3) heat inactivated fetal bovine serum (Gemini Bioproducts) and incubated at 37°C in 5% CO₂.

Protein Assays. Cells were grown to 60-70% confluence and exposed to drugs or DMSO vehicle for the indicated time periods. Lysates were prepared using 50 mM Tris pH 7.4, 2% SDS and 10% glycerol lysis buffer. Protein concentration was determined using the BCA kit (Pierce Chemical Co.), according to the manufacturers instructions. Clarified protein lysates (20-50 mg) were electrophoretically resolved on denaturing SDS-PAGE, transferred to nitrocellulose and probed with the following primary antibodies: anti-Her2 (C-18), -Her3 (C-17), -Raf-1, -cyclin D1, -Rb (C-15) (Santa Cruz Biotechnology), anti-hsp90, -hsp70 (Stressgen) anti-Trap-1 (MSK81), anti-b-actin, -tubulin (Sigma), ER, anti-PI3K (p85) (Upstate Biotechnologies).

Antiproliferative index. Growth assays were performed by seeding 10000 cells (MCF-7 and MDA-MB-468) and 20000 cells (SKBr3) per well in 6-well dishes and incubating for 24 hrs before drug treatment. Drugs or vehicle were administered as outlined for each experiment, and cells were incubated for the time periods depicted and then the number quantified by a coulter counter.

Tissue Culture IC₅₀ Studies. Growth inhibition studies were performed using the sulforhodamine B assay described before. Experiments were performed with BT-474, MDA-MB-468, MCF-7, and TSU-Pr1. Stock cultures were grown in T-175's flask containing 30 mL of DME (HG, F-12, non-essential amino acids, and penicillin and streptomycin), with glutamine, and 10% FBS. TSU-Pr1 were grown in RPMI 1640 with glutamine and 10 % FBS. Cells were dissociated with 0.05% trypsin and 0.02% EDTA in PBS without calcium and magnesium.

Experimental cultures were plated in microtiter plates (Nunc) in 100 uL of growth medium at densities of 1000 cells per well, except for BT-474 which were plated at densities of 3000 cells per well. One column of wells was left without cells to serve as the blank control. Cells were allowed to attach overnight (BT-474 were allowed to attach for 48 hours). The following day, an additional 100 uL of growth medium was added to each well. Stock drug or DMSO was dissolved in growth medium at twice the desired initial concentration. Drug or DMSO was serially diluted at a 1 : 1 ratio in the microtiter plate

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and added to duplicate wells. After 72 hours of growth, the cell number in treated versus control wells was estimated after treatment with 10% trichloroacetic acid and staining with 0.4% sulforhodamine B in 1% acetic acid. The IC_{50} is calculated as the drug concentration that inhibits cell growth by 50% compared with control growth.

Her2 degradation. Total Protein Assays. Cells were grown to 60-70% confluence and exposed to drugs or DMSO vehicle for the indicated time periods. Lysates were prepared using 50 mM Tris pH 7.4, 2% SDS and 10% glycerol lysis buffer. Protein concentration was determined using the BCA kit (Pierce Chemical Co.), according to the manufacturers instructions. Clarified protein lysates (20-50 μ g) were electrophoretically resolved on denaturing SDS-PAGE, transferred to nitrocellulose and probed with the anti-Her2 primary antibody (C-18) (Santa Cruz Biotechnology).

Binding studies. Solid phase competition assays. GM was immobilized on Affigel 10 resin (BioRad) as described². The GM-beads were washed with TEN buffer (50 mM Tris:HCl pH 7.4, 1 mM EDTA, 1% NP-40) containing protease inhibitors and then blocked for 45 min at 4 °C with 0.5% BSA in TEN buffer. Hsp90 protein from Stressgen (SPP-770) was incubated with or without drugs for 17 min on ice. To each sample were added 20 μ L GM-beads and the mixtures were rotated at 4 °C for 1 hr followed by 3 washes with 500 μ L ice cold TEN each. The GM-beads bound protein was eluted from the solid phase by heating in 65 μ L 1xSDS. Samples were portioned in a 20 μ L aliquot for Hsp90 alpha analysis and a 40 μ L aliquot for Hsp90 beta analysis, applied to a SDS/PAGE gel and visualized by immunoblotting with Hsp90 alpha (Stressgen # SPA-840) and Hsp90 beta (NeoMarkers#RB-118), respectively.

Compounds PU4-72 were tested for binding to Hsp90, degradation of Her2 total protein and for their antiproliferative effect. The results are summarized in Tables 1, 2, 3 and 4.

Table 1. Influence of the nature of the 9-N chain on activity

EC_{50}	EC_{50}	IC_{50}	IC_{50}	IC_{50}	IC_{50}
Hsp90 α	Hsp90 β	MCF-7	Her2/MC F-7	BT-474	MDA-468

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PU13	4			70		75
PU22	5			80		118
PU43	6	6.6	10.8	47	50	54
PU3	7	15	13	50	55	61
PU21	8	22.5	11.8	62	50	73
PU41	9	16.6	20.4	98	100	70
PU9	10	28	114	69		85
PU14	11	62.3	52.1	160	>100	130
PU26	13	25.3	15.7	62	70	71
PU15	15	12	16.2	75	70	70
PU30	16	111.3	47.8	111	120	
PU16	17	17	32	46	60	68
PU4	18	120	>100	70	50	
PU23	22	13.3	9.5	47	50	55
PU7	23	17.6	7.5	64	70	80
PU8	24	4	10.6	41	30	73
PU11	25			51		66
PU24	26	2.6	1.5	24	20	41
PU25	36			82		
PU44	37	4.1	12.4	65	85	79
PU29	38	1.4	1.7	39	45	72
PU20	41	4.4	3.4	92	70	80

*all other compounds from Fig. 6B were either inactive or insoluble

Table 2. The influence of C-2 fluorination on activity

		EC ₅₀	EC ₅₀	IC ₅₀	IC ₅₀	IC ₅₀	IC ₅₀
		Hsp90 α	Hsp90 β	MCF-7	Her2/MC F-7	BT-474	MDA-468
PU43F	47	4.1	5.3	25	30	30	25
PU3F	48	6	3.5	24	25	29	30
PU21F	49	22	8.9	36	35	43	19
PU44F	50	7.9	13.8	64	65	60	35
PU26F	51	22	15	44	40	45	28
PU29F	52	2	1.3	16	15		16
PU8F	53	5	9.8	33	30	35	25
PU47F	54	10.7	9.9	45	50	46	33
PU48F	55	10	3.5	25	20	28	16
PU49F	56	15	10.4	37	45	33	45
PU24F	57	6.2	0.7	11	5	14	15
PU16F	58	30.7	9.2	41	40	31	21
PU20F	59	2.3	6.8	23	20	25	21

Addition of either CN, vinyl, iodine, methoxy, ethoxy, NH₂ at position 2 of the purine moiety decreased or abolished activity.

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Table 3. The influence of introduction of an electron-donor group on the phenyl moiety

		EC ₅₀	EC ₅₀	IC ₅₀	IC ₅₀	IC ₅₀	IC ₅₀
		Hsp90 α	Hsp90 β	MCF-7	Her2/MC F-7	BT-474	MDA-468
PU3	7	15	13	50	55		61
PU3PhCl	65	18.6	4.6	19	25	30	36
PU3PhBr	66	35.3	11.7	25	50	60	INSOL
PU3PhBr ₂	67	>100	>100	30	>100		21

Assimilation of the best substituents resulted in the derivatives 71 and 72 (Table 4).

Table 4.

PU24FCI	71	0.55	0.45	2	2	4.5	3
PU29FCI	72	0.65	0.52	5.4	3	4.5	4.5

EXAMPLE 9

The ability of PU3 and PU24FCI to induce degradation of Akt protein. As shown in Figs. 17 A and B, the 30-fold difference in activity which was observed for Her2 degradation was also reflected in degradation of Akt. An approximately 30-fold difference was also seen in the Hsp90 binding affinity.

EXAMPLE 10

Data was analyzed to see if there was a correlation between concentrations of PU-series compounds effective for inducing Her2 degradation and the concentrations required for growth inhibition in the MCF-7 cell line, and between the Hsp90 binding constant and the concentrations required for growth inhibition. As shown in Figs 18 A and B, the experimental results suggest a good correlation between growth arrest, Her2 total protein degradation and Hsp90 binding efficacy.

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EXAMPLE 11

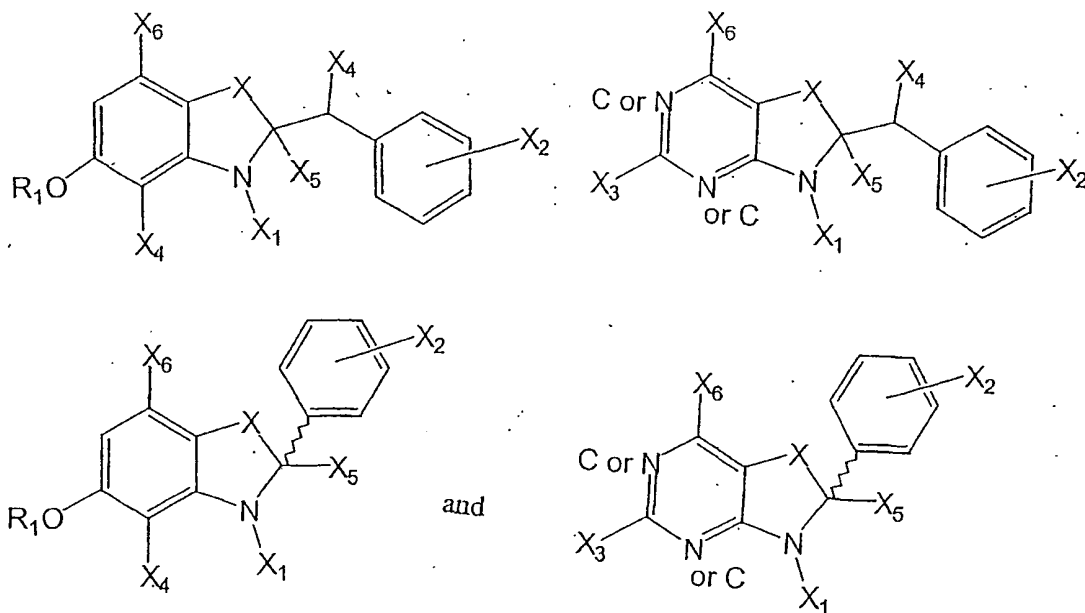
Cell culture. The human cancer cell lines MCF-7, SKBr3 and BT-474 were obtained from the American Type Culture Collection (Manassas, VA) and maintained in 1:1 mixture of DME:F12 supplemented with 2 mM glutamine, 50 units/mL penicillin, 50 units/mL streptomycin and 5% (for MCF-7) or 10% (for SKBr3 and BT-474) heat inactivated fetal bovine serum (Gemini Bioproducts) and incubated at 37°C in 5% CO₂.

Her2 degradation. Total Protein Assays. Cells were grown to 60-70% confluence and exposed to drugs or DMSO vehicle for the indicated time periods. Lysates were prepared using 50 mM Tris pH 7.4, 2% SDS and 10% glycerol lysis buffer. Protein concentration was determined using the BCA kit (Pierce Chemical Co.), according to the manufacturers instructions. Clarified protein lysates (20-50 µg) were electrophoretically resolved on denaturing SDS-PAGE, transferred to nitrocellulose and probed with the anti-Her2 primary antibody (C-18) (Santa Cruz Biotechnology). As shown in Fig. 19, PU24FC1 induces efficient degradation of the oncogenic protein Her2 at low concentrations (IC₅₀ of 1.7, 2, 4.5µM for SKBr3, MCF-7 and BT-474 respectively).

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What is claimed is:

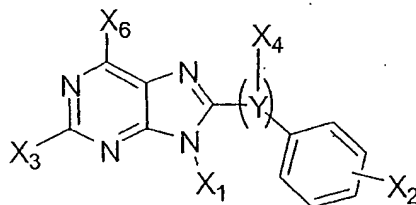
1. A compound comprising a binding moiety which binds to the N-terminal pocket of a member of the HSP90 family of proteins, wherein the compound is a synthetic molecule and wherein the binding moiety is not an ansamycin antibiotic or radicicol or a derivative thereof.
2. The compound in accordance with claim 1, wherein the compound is soluble in aqueous solution.
3. The compound of claim 1, wherein the binding moiety is a substituted purine.
4. The compound of claim 1, wherein the binding moiety is coupled to a targeting moiety which specifically binds to a protein receptor or marker.
5. The compound according to claim 1, wherein the binding moiety is selected from the group consisting of:



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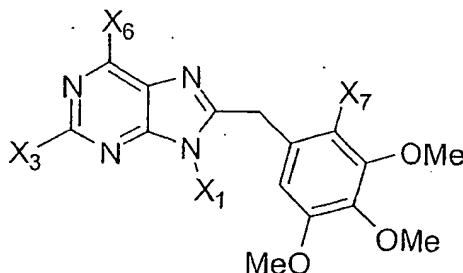
wherein X is O, S, NH, CH₂, or CO; X₁ is a hydrophobic chain or COR, wherein R is a hydrophobic chain; X₂ is from 1 to 5 hydrogen acceptor/donor functionalities, which may be the same or different; X₃ is a small size group selected from among alkyl groups (saturated, unsaturated, cyclic or linear), alkoxy groups, halogen, SMe, and SEt; X₄ is H or as X₃; X₅ is H or as X₁; X₆ is -NH₂, -OH, -OAlkyl, -CONH₂ or a similar hydrogen donor functionality and R₁ is H or alkyl.

6. The compound according to claim 1, wherein the binding moiety has the formula:



wherein Y is C, O, N, or O-C; X₁ is a hydrophobic chain or COR, wherein R is a hydrophobic chain; X₂ is from 1 to 5 hydrogen acceptor/donor functionalities, which may be the same or different; X₃ is a small size group selected from among alkyl groups (saturated, unsaturated, cyclic or linear), alkoxy groups, halogen, SMe, and SEt; X₄ is absent, H or as X₃; X₅ is H or as X₁; X₆ is -NH₂, -OH, -OAlkyl, -CONH₂ or a similar hydrogen donor functionality and R₁ is H or alkyl.

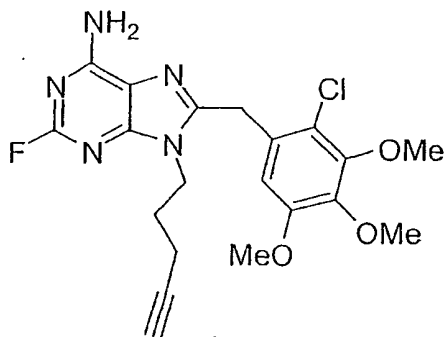
7. The compound of claim 6, wherein the binding moiety has the structure



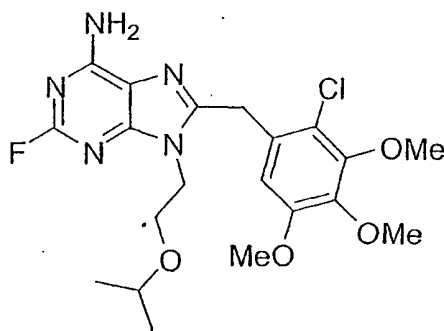
- 34 -

wherein X₇ is hydrogen or halogen and X₃ is halogen.

8. The compound according to claim 7, wherein the binding moiety has the structure



9. The compound according to claim 7, wherein the binding moiety has the structure



10. The composition according to any of claims 1-9, wherein a first binding moiety is coupled to a second binding moiety to form a dimer.

11. A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a compound comprising a binding moiety which binds to the N-terminal pocket of a member of the HSP90 family of proteins, said binding moiety having a backbone which can achieve a folded C-configuration when disposed within the N-terminal pocket of a member of the HSP90 family of proteins, and substituents on the backbone directed in orientations to interact with a plurality of potential binding sites in the N-terminal pocket to an extent that the binding moiety has a binding affinity for the

- 35 -

N-terminal pocket that is greater than ADP and different from geldanamycin for at least one member of the HSP90 family of proteins, and provides differential degradation of proteins requiring chaperones of the HSP90 family.

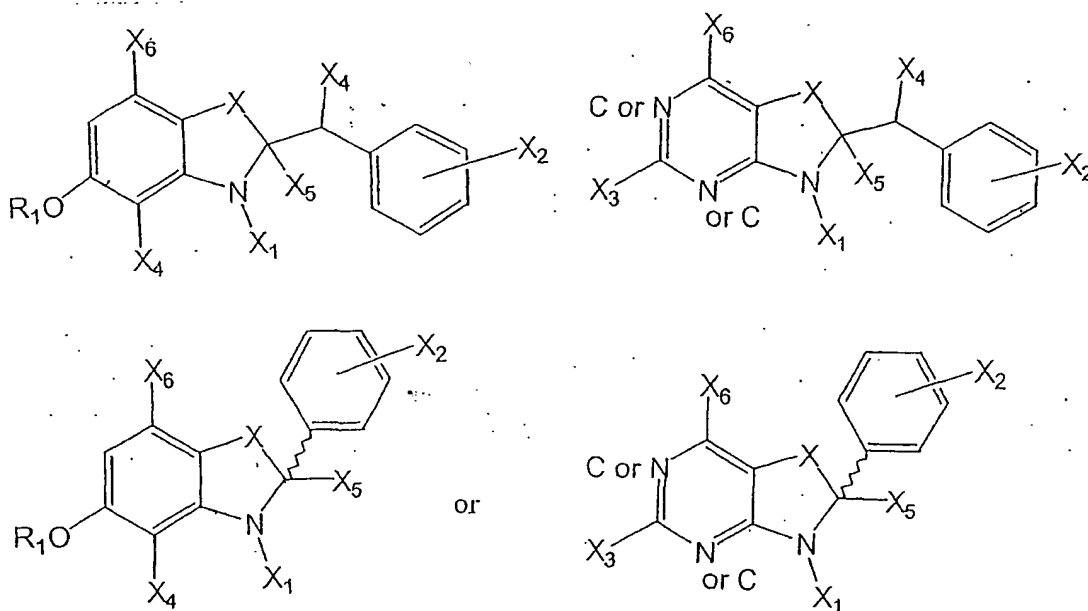
12. The composition according to claim 11, wherein the binding moiety is part of or all of a compound in accordance with any of claims 1-8.

13. The composition according to claim 11, wherein the binding moiety is part of or all of a compound in accordance with claim 9.

14. A method for inhibiting the growth of cells which are dependent on a protein or receptor requiring a member of the HSP90 family for function, comprising the step of treating the cells with a composition according to any of claims 1-8.

15. A method for inhibiting the growth of cells which are dependent on a protein or receptor requiring a member of the HSP90 family for function, comprising the step of treating the cells with a composition according to claim 9.

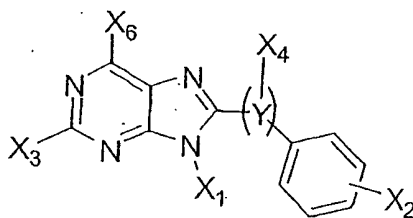
16. A compound having the formula:



- 36 -

wherein X is O, S, NH, CH₂, or CO; X₁ is a hydrophobic chain or COR, wherein R is a hydrophobic chain; X₂ is from 1 to 5 hydrogen acceptor/donor functionalities, which may be the same or different; X₃ is a small size group selected from among alkyl groups (saturated, unsaturated, cyclic or linear), alkoxy groups, halogen, SMe, and SEt; X₄ is H or as X₃, X₅ is H or as X₁; X₆ is -NH₂, -OH, -OAlkyl, -CONH₂ or a similar hydrogen donor functionality and R₁ is H or alkyl.

17. A compound having the formula:



wherein Y is C, O, N, or O-C; X₁ is a hydrophobic chain or COR, wherein R is a hydrophobic chain; X₂ is from 1 to 5 hydrogen acceptor/donor functionalities, which may be the same or different; X₃ is a small size group selected from among alkyl groups (saturated, unsaturated, cyclic or linear), alkoxy groups, halogen, SMe, and SEt; X₄ is absent, H or as X₃, X₅ is H or as X₁; X₆ is -NH₂, -OH, -OAlkyl, -CONH₂ or a similar hydrogen donor functionality and R₁ is H or alkyl.

18. A method for making an 8-benzyl purine composition comprising the steps of:

- (a) converting a carboxylic acid starting material to an acid halide; and
- (b) reacting the acid halide with an amino-substituted pyrimidine to produce an intermediate product which undergoes ring closure an 8-benzyl purine.

19. The method of claim 18, further comprising the step of alkylating the 8-benzyl purine at the nitrogen at the 9-position.

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20. The method of claim 19, wherein the alkylation is performed using a Mitsunobu alkylation reaction.

21. A method for screening a test compound for binding affinity to the N-terminal pocket of a member of the HSP90 family, comprising the steps of:

- (a) contacting the test compound and the member of the HSP90 family with a substrate having an hsp binding moiety immobilized thereon; and
- (b) observing the amount of the member of the HSP90 family bound by the immobilized ansamycin antibiotic, wherein a reduction in the amount of the member of the HSP90 bound as compared to the amount bound in the absence of the test compound indicates that the test compound binds to the member of the HSP90 family.

22. The method according to claim 21, wherein the hsp binding moiety is an ansamycin antibiotic or radicicol.

23. The method according to claim 21, wherein the hsp binding moiety is geldanamycin.

24. The method according to claim 21, wherein the hsp binding moiety is a compound in accordance with any of claims 1-9, 16 or 17.

25. A method for screening a compound for selective binding to Hsp90-alpha or beta, comprising the steps of:

- (a) combining a support having a non-specific HSP90-binding moiety with a solution containing an Hsp90 preparation that includes both Hsp90-alpha and beta in the presence of the compound to be screened;
- (b) detecting the amount of Hsp90-alpha and beta that bind to the support, wherein a compound with differential binding selectivity will inhibit binding of Hsp90-alpha and beta to the support to different extents.

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26. The method of claim 25, wherein the non-specific HSP90 binding moiety is geldanamycin.

21 * AFQAEIAQLM SLINTEYSN KEPLRELISS NSSDALDKIR
81 ⁵ EVNRNM KLINSLYKN KEPLRELISS NASDALDKIR LISL
21 # AOLM SLINTEYSN KEPLRELISS NASDALDKIR YESLT
91 @ QABTKKLLDIVARSLYSE KEVFIREFLISS NASDALEKIR HK.

61 * YETLTDPSKL DSGKELHNL IPNKQDRILT IVDTGIGMTK ADLNNLGTI AKSGTKAFME
121 ³ TDENALSGNE ELTVKIKCDK EKNLHVTDITGYGMTR BELVKNLGTI AKSGTSEFLNKMTE
61 # DPSKLDGKE LKIDIPNPQ ERTLT LVDTGIGMTK ADLNNLGTI AKSGTKAFME
131 @ LVSDGQALPE MEHLQTNAE KGTITIQTIGMTQ EELVSNLGTI ARSGSKAFDLALQNQ

121 * ALQAGADISM IGQFGVGFYS AYLVAEKVTV ITKHNDDEQ YAWESSAGGSF TVRTDTGPM
181 ⁵ AQEDGOSTSEL IGQFGVGFYS A FLVADKVTV ITKHNDIQ HIWESDSNEF SVIADPRGNT
116 # ALQAGADI SM IGQFGVGFYS AYLVAEKVVV IRKHNDDEQ YAWESS AGGSFTVRAD HGEPIGM
186 @ ALQNOABASSKI IGQFGVGFYS AFMVADRVVYSR SAAPGSLGYQ WLSDGSGVFE

181 * GRGTVKVLH LKEDQTEYLEE RRIKHVKKH SQFIGYPITL FVEKERDKEV SDDEAEEKED
241 ⁵ LGRGTTITLV LKEEASDYLE LDTKNLVKK YSQFNFPY VWSSKTETVE EPMEEEA
176 # GMGTVKVLH LKEDQTEYLEE RRVKEVVKKHQFIG
246 @ GVRITGKII I H LKS DKEFSSEARV RDVVVKYSNF

* HSP90 ALPHA (P07900); ⁵ GRP94 (NP_003290); # HSP90 BETA (P08238); @ TRAP1 (AAF15314)

Fig 1

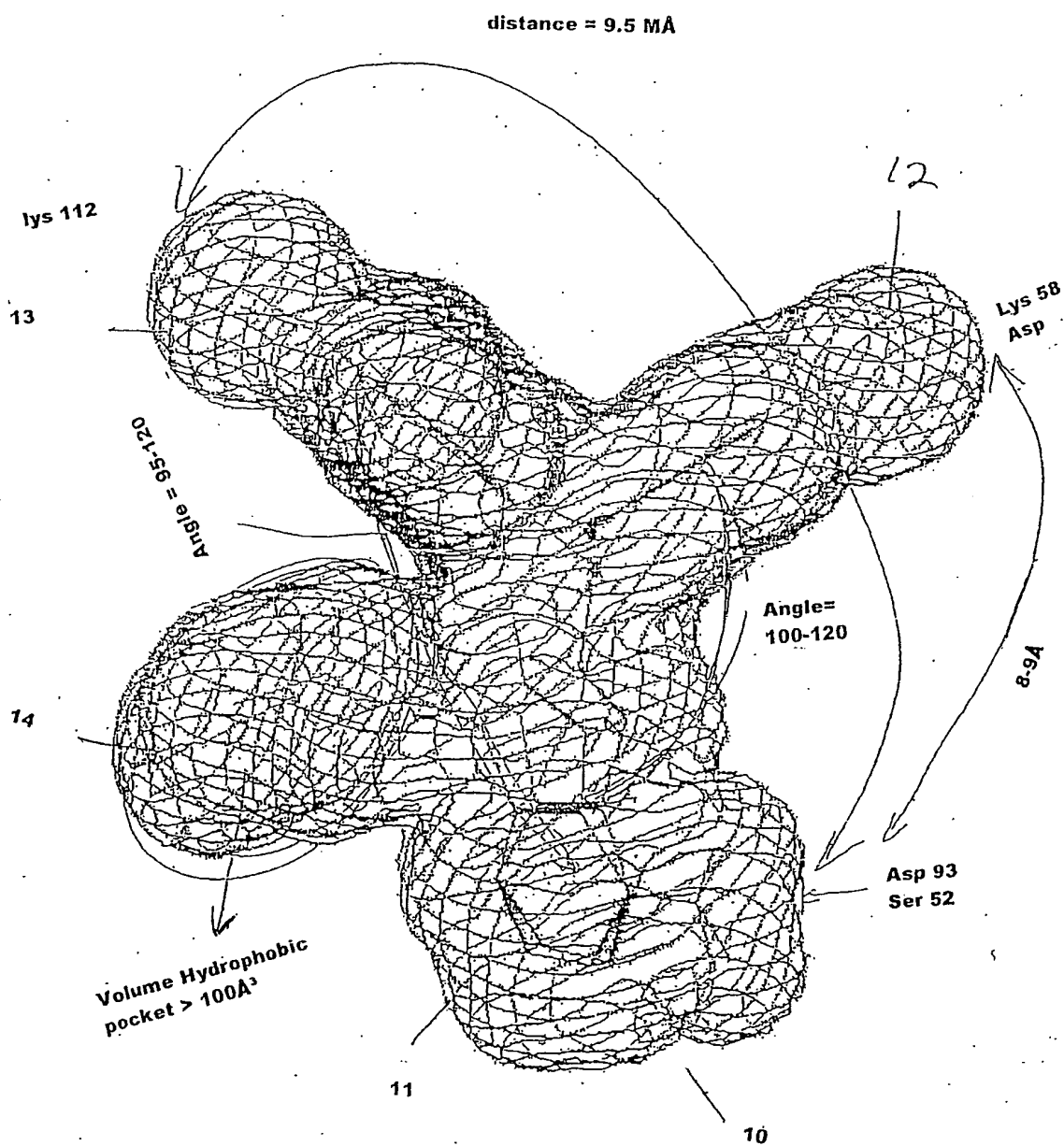


Fig 2

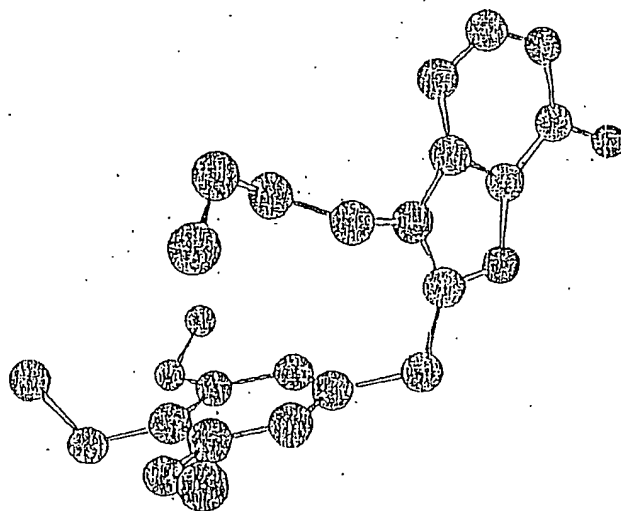


Fig 3B

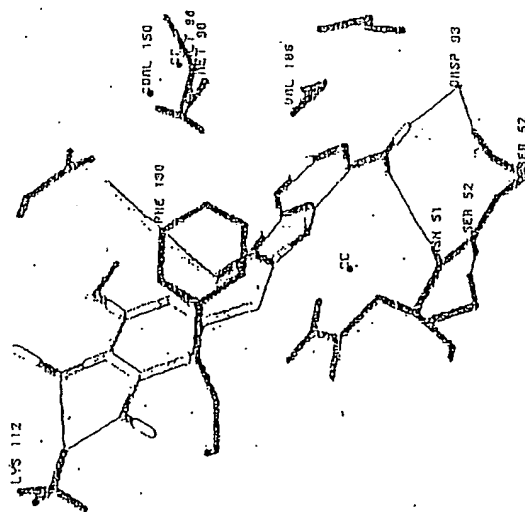
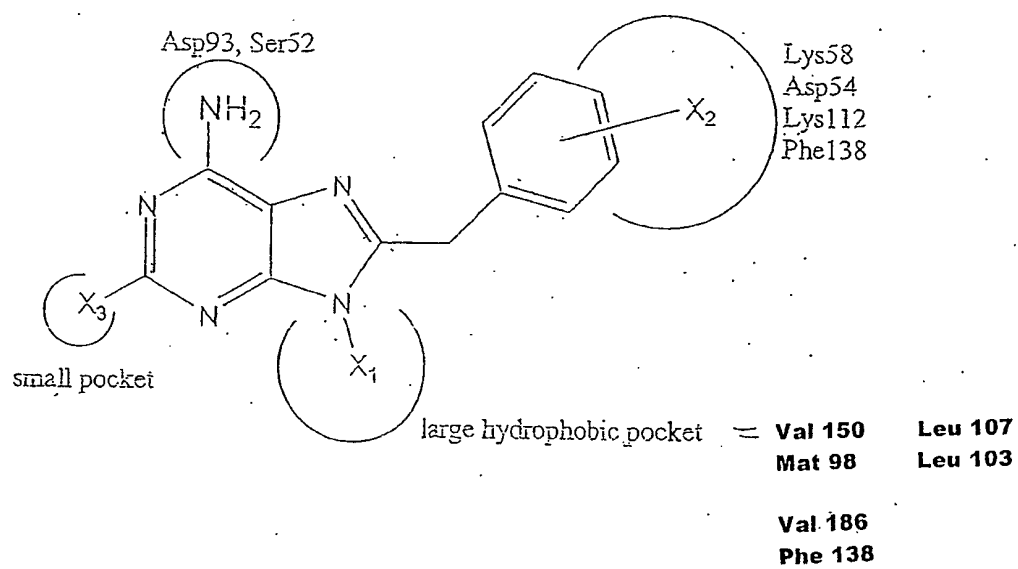


Fig 3A

**Fig 4**

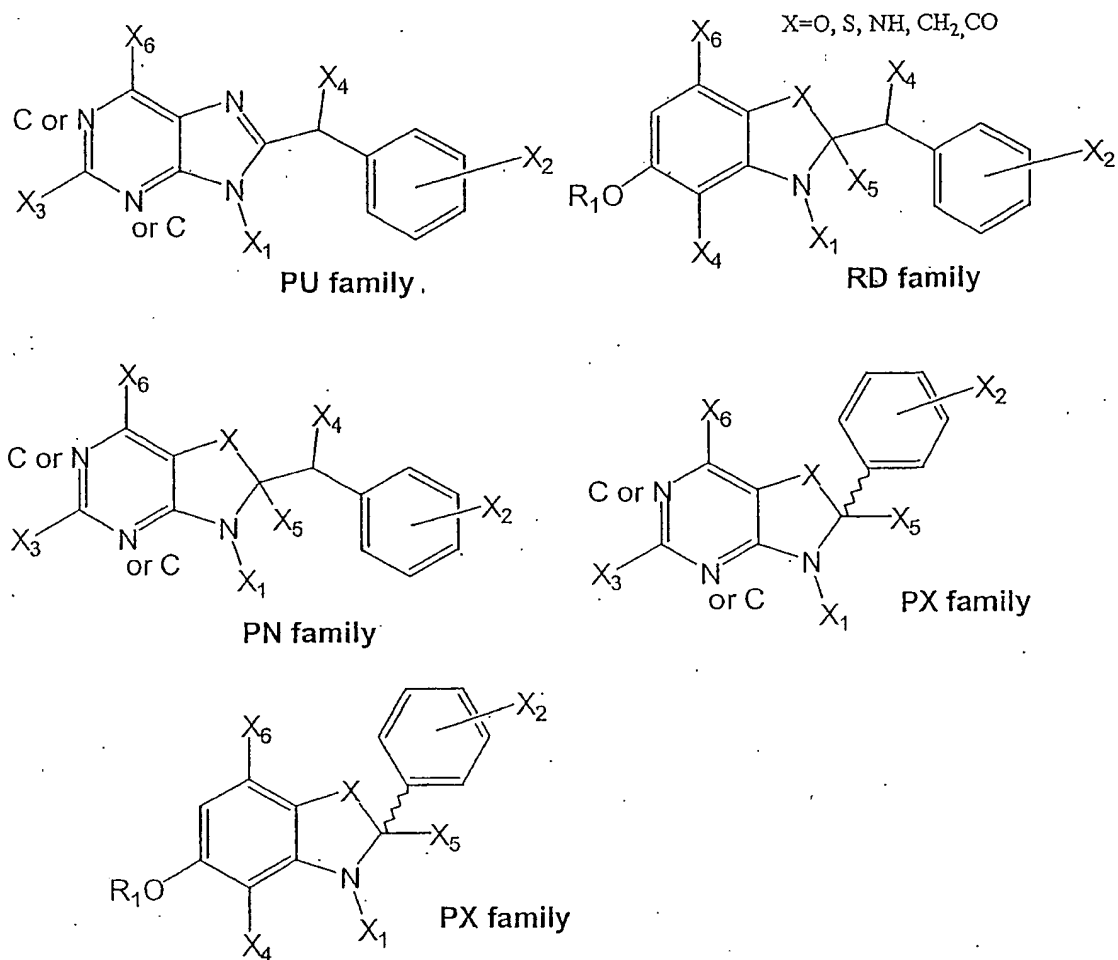


Figure 5

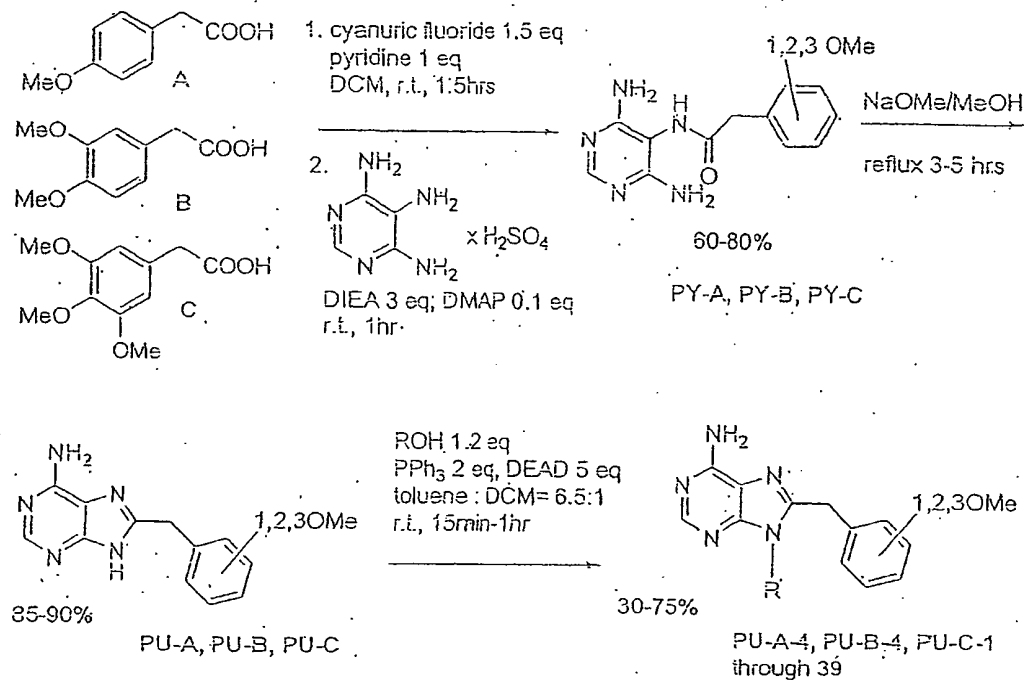


Fig 6A

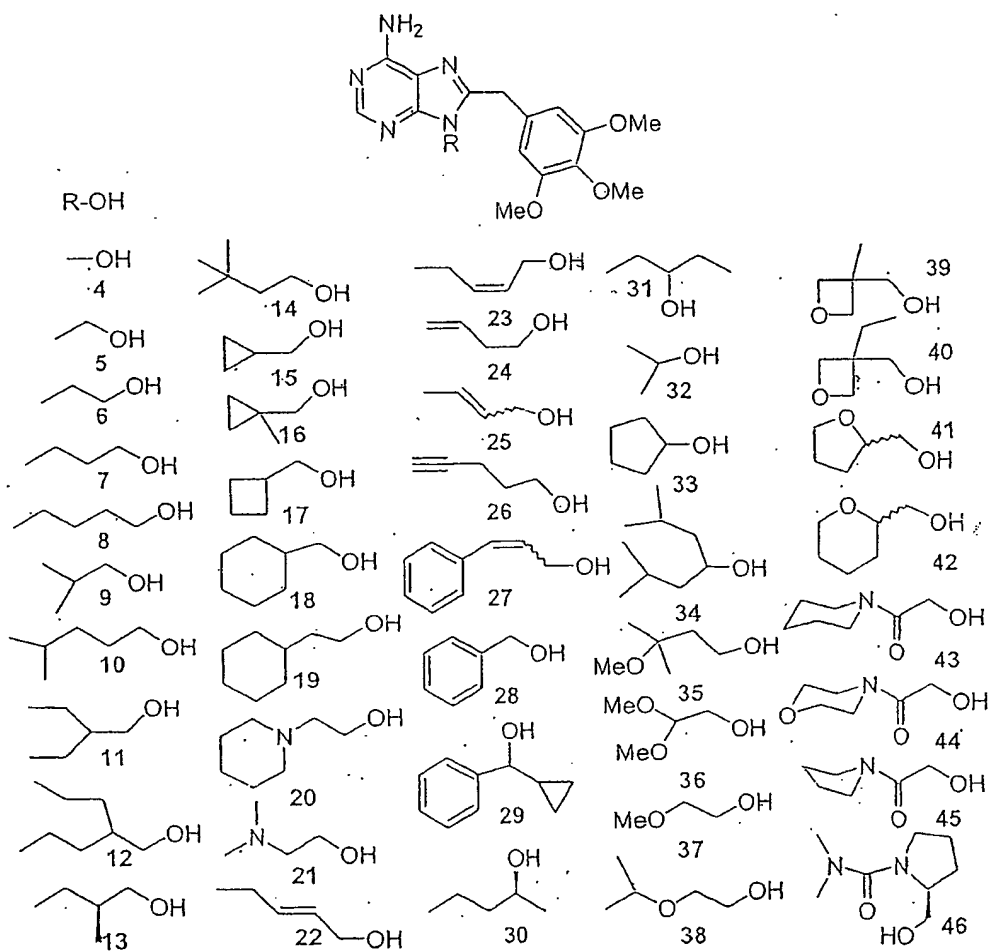


Fig 6B

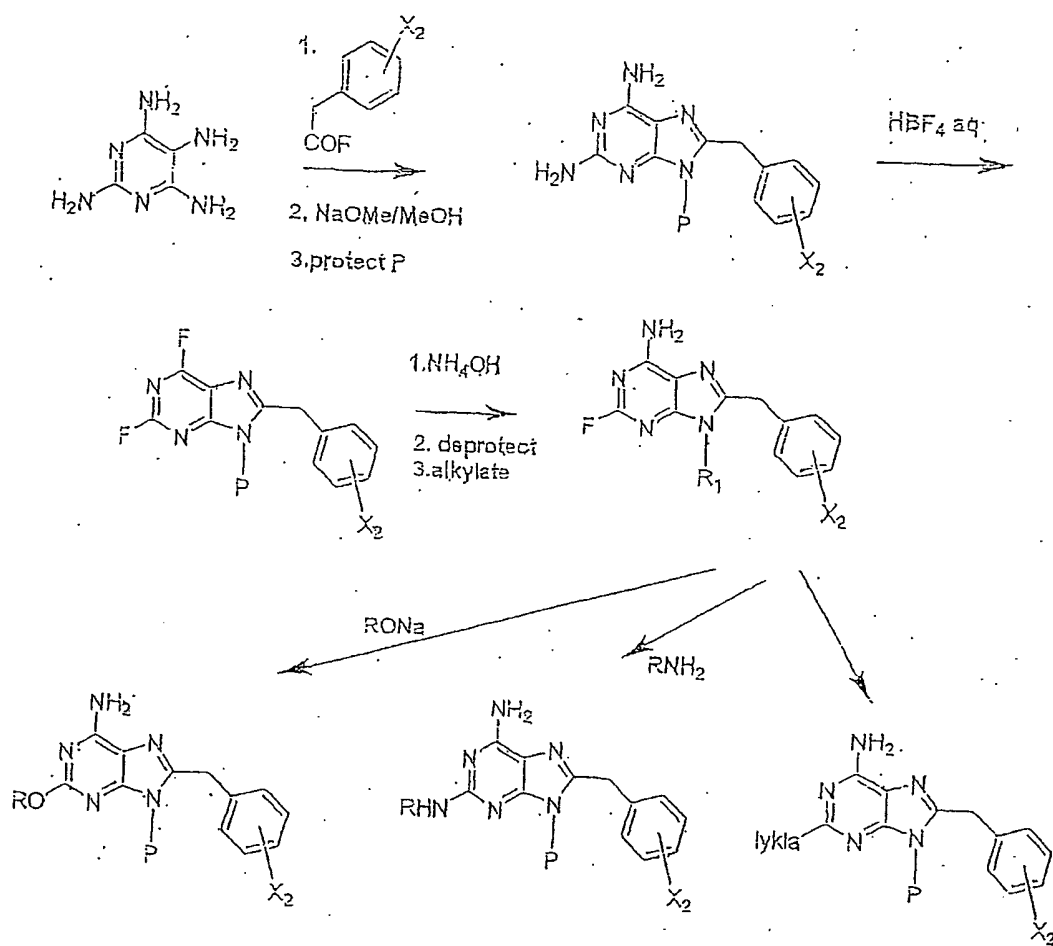
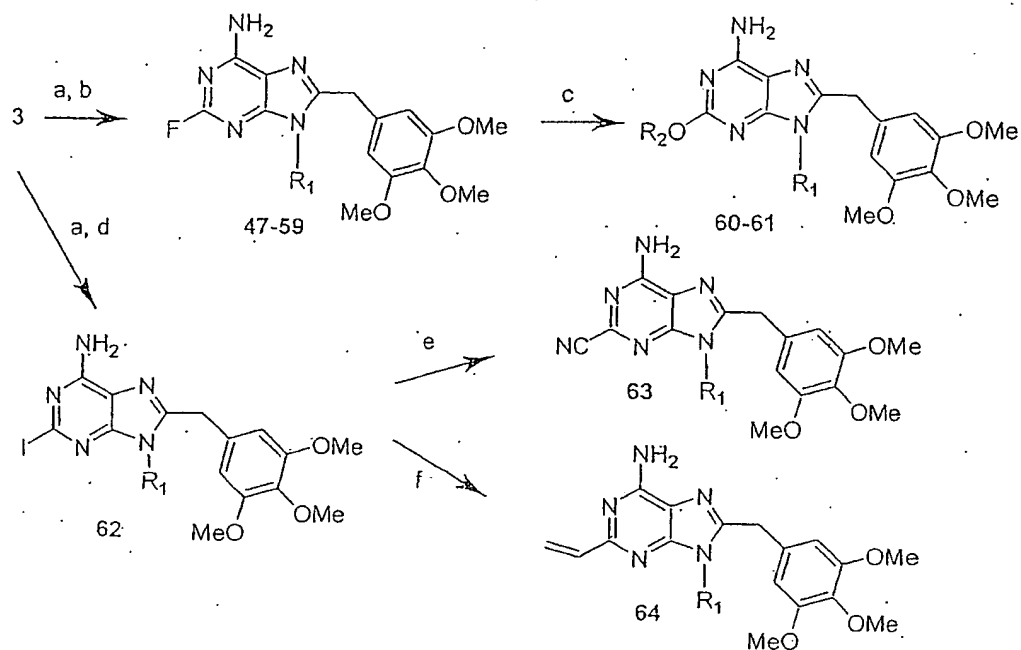


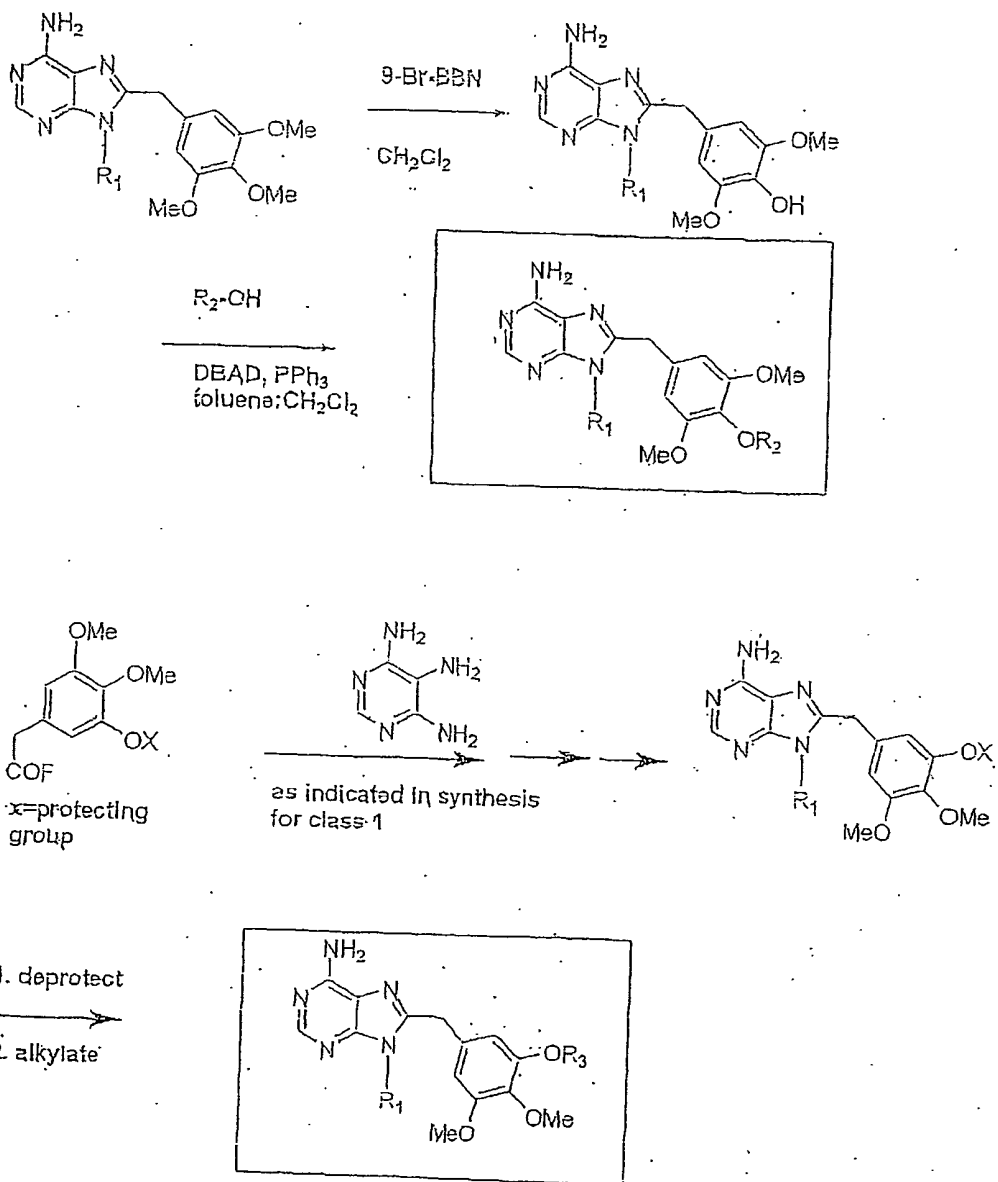
Fig 7A



Reagents: (a) HF, pyridine, *t*-butyl nitrite; (b) R₁OH, PPh₃, diethyl azodicarboxylate or di-*tert*-butyl azodicarboxylate, toluene:DCM; (c) R₂OH, NaOMe; (d) CH₂I₂, isoamyl nitrite; (e) Bu₃SnCN, Pd[PPh₃]₄; (f) (MeCN)₂PdCl₂, Bu₃(vinyl)Sn.

Fig 7B

Synthesis of class II small molecules. Variations of X2.



R2 and R3 any

Fig 8A

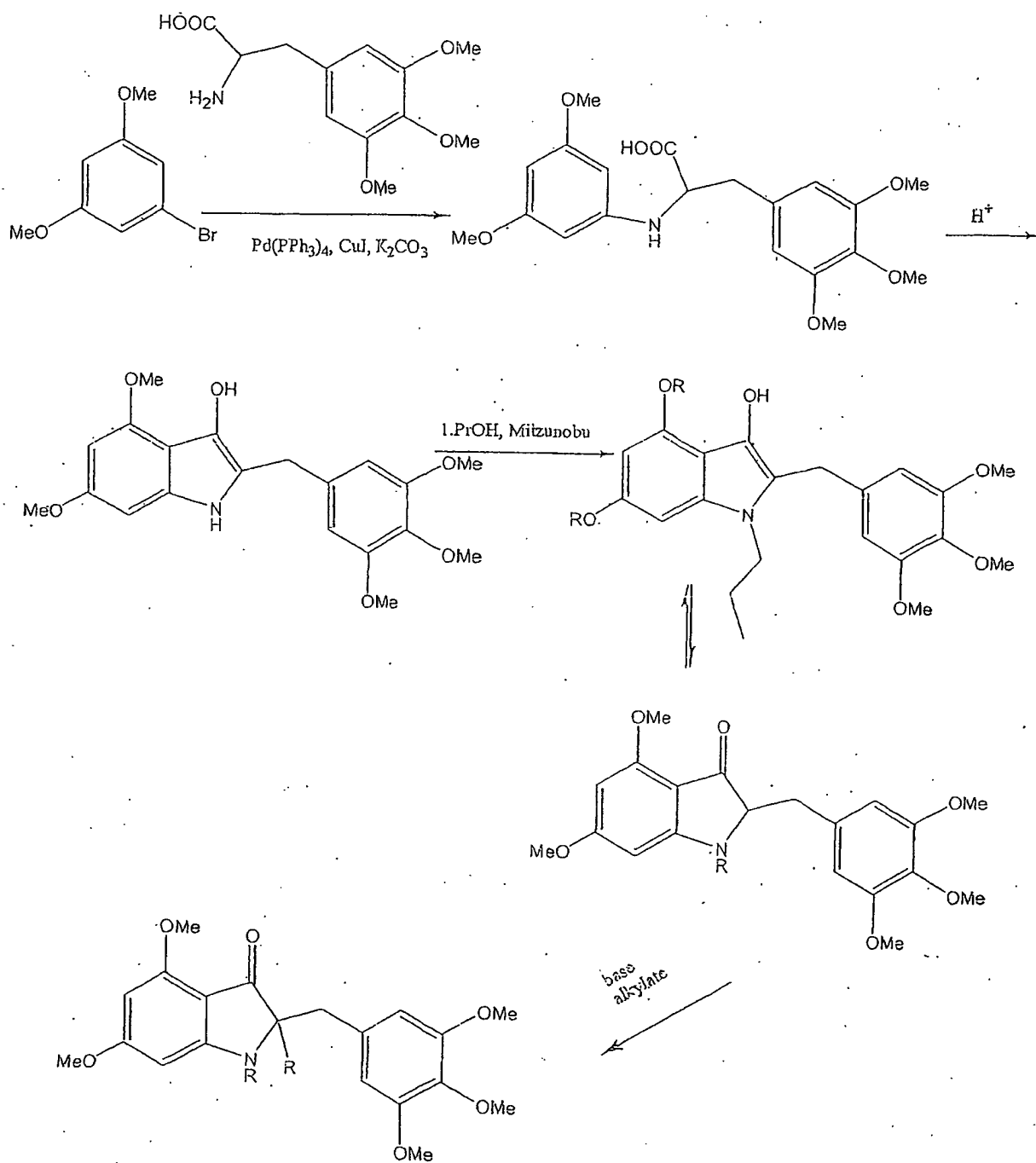
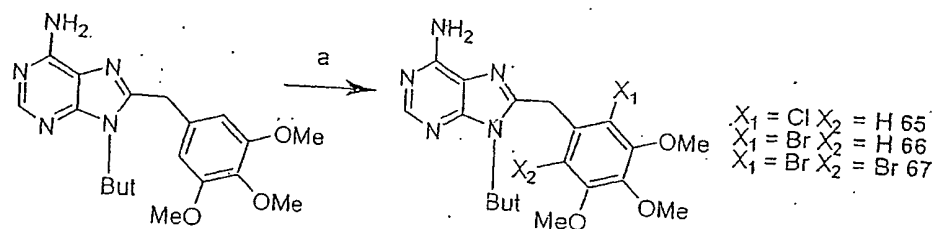
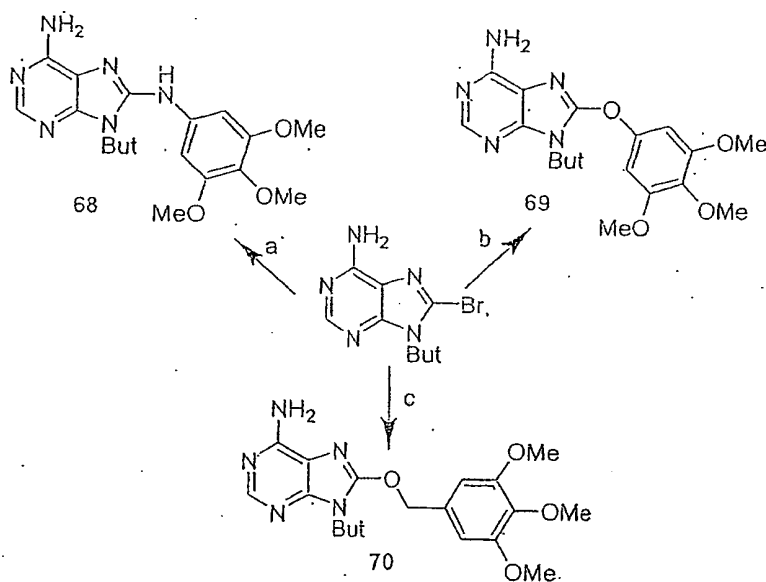
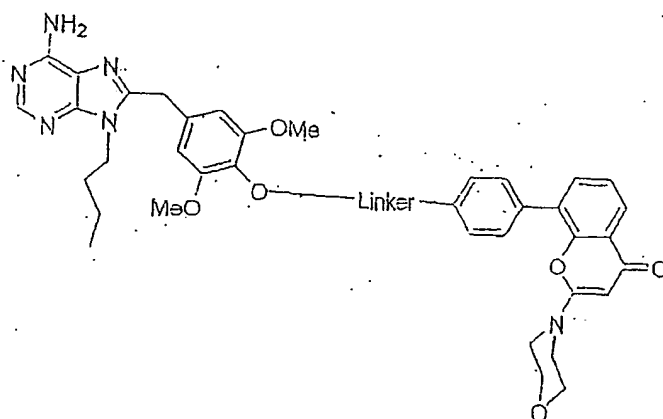
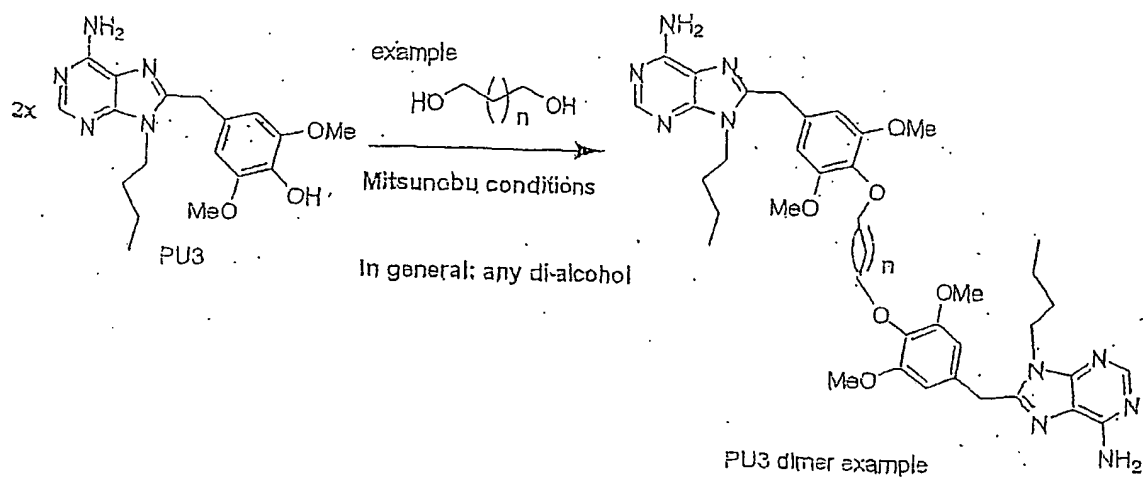


Fig 9

**Fig 8B****Fig 10**

Mode of attachment of PU3 (as example molecule of PU family) to other entities [another PU3=dimer, an inhibitor or a carrier molecule=antibody etc]



General example of PU3 heterodimer:
PU3 linked to the PI3K inhibitor, LY294002

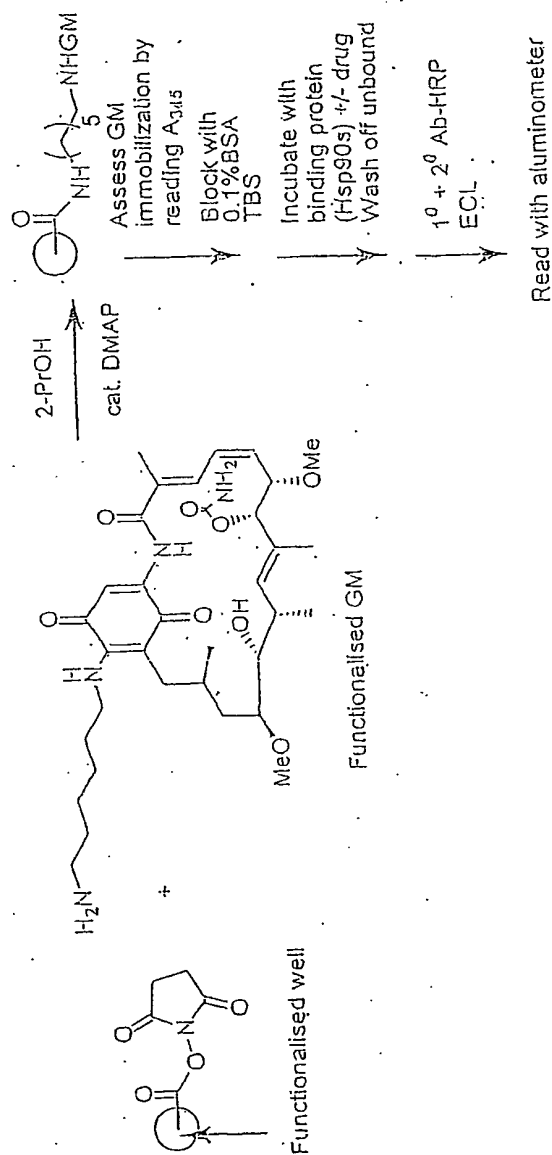


Fig 12

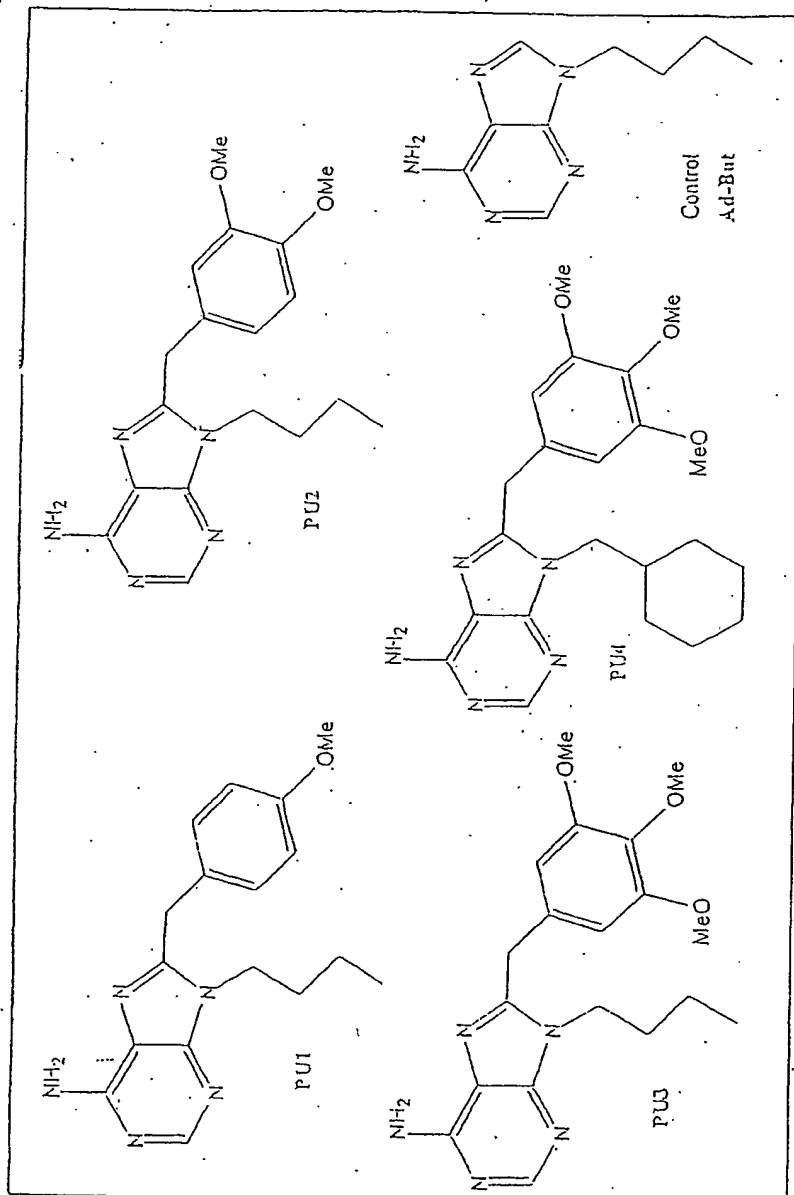


Fig 13

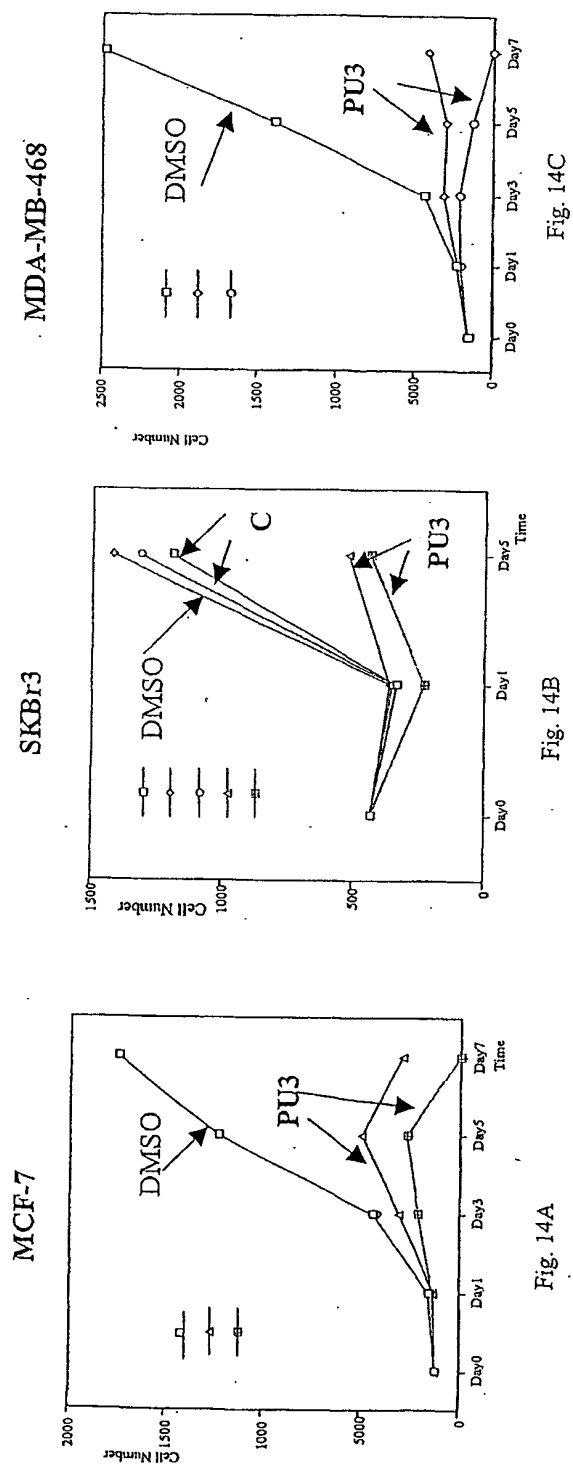


Fig. 14C

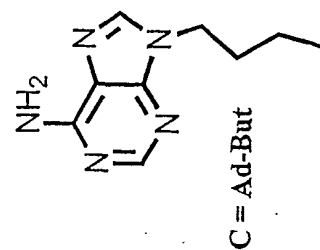
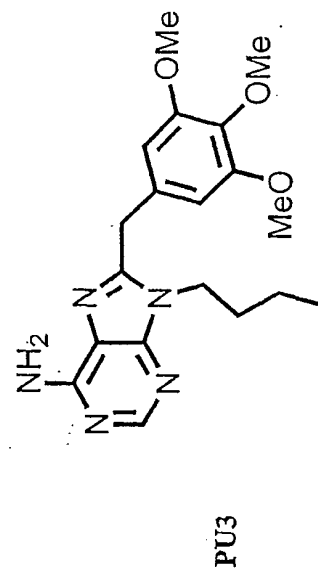
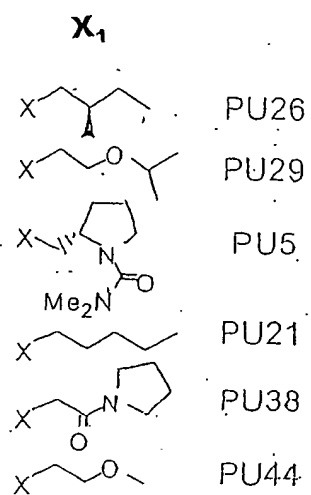
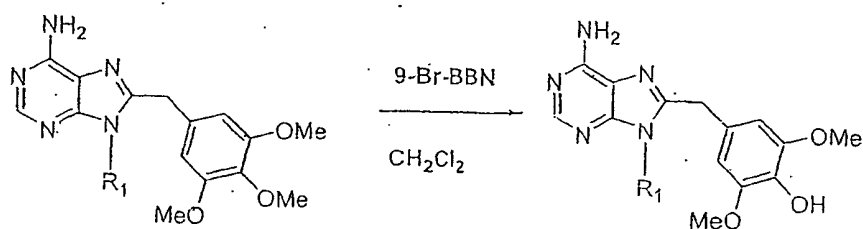


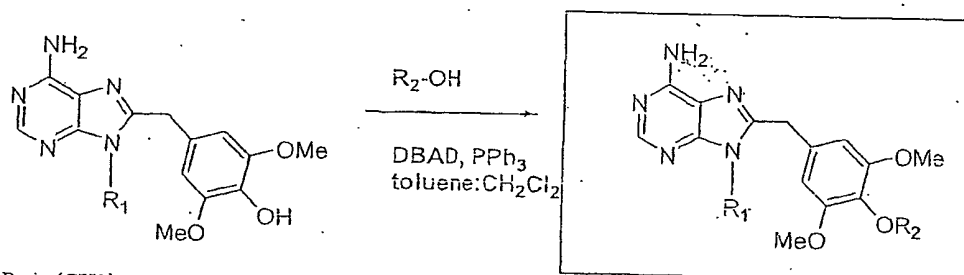
Fig. 14B



**Fig 15**



Charge 50mg of PU3 to 4mL anhydrous CH_2Cl_2 . Charge 2mL of 1M 9-BrBBN in CH_2Cl_2 (15equivalents). Stir for 1hr at room temperature. Add 4mL NaHCO_3 . Stir 15min. Split and extract aqueous with 3x5mL CH_2Cl_2 . Concentrate. Column on silica gel with 10:5:5:1.5=hexanes: CH_2Cl_2 :EtOAc:MeOH. 25mg product=52% yield.

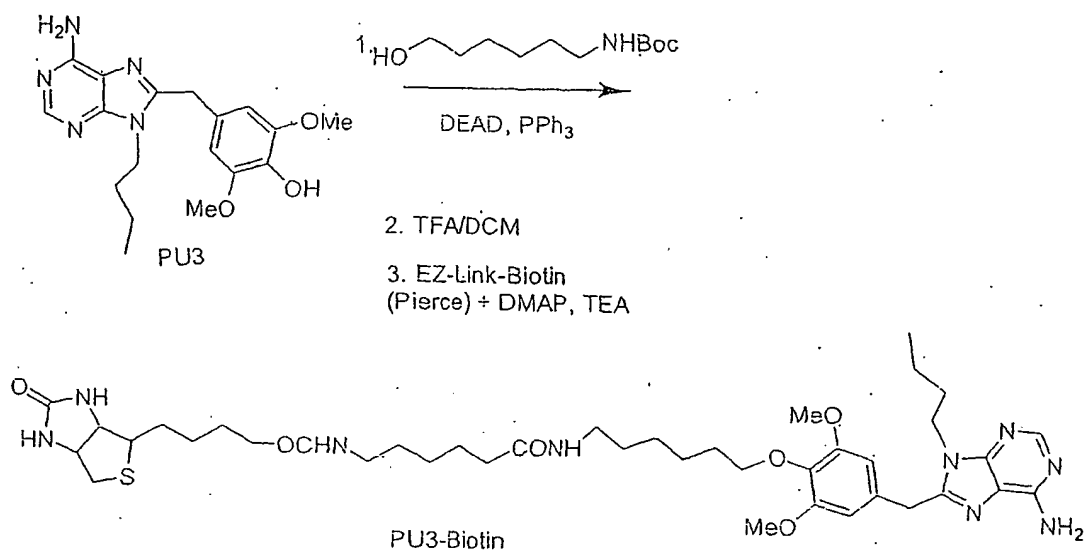


R_1 is $(\text{CH}_3)_2\text{CHCH}_2\text{-}$

5mg of SM (0.014mmol), 8mg PPh_3 (2eq), 16mg di-*t*-butylazodicarboxylate (5equivalents), 2 μL ROH (1.5eq) in 0.4mL CH_2Cl_2 were added till all dissolved. Add 1.3mL toluene. Stir at RT 16hrs.. Concentrate. Column silica gel with 10:5:5:1.5=hexanes: CH_2Cl_2 :EtOAc:MeOH.

ROH	yield%
	17
	43
	35

Fig 16A



Exp. To Sepharose-Streptavidin [40uL/exp] were added increasing concentrations of PU3-Biotin and the slurry equilibrated for 15min at rt. Beads were washed with PBS 0.2% Tween and 30uL of RSBT buffer were added to each sample, followed by identical amounts of proteins [Hsp90 or Trap]. Samples were rocked for 1hr at 4°C. After several washes the amount of bound protein was eluted by boiling in 50uL 1xSDS buffer. Proteins were assessed by immunoblotting.

Fig 16B

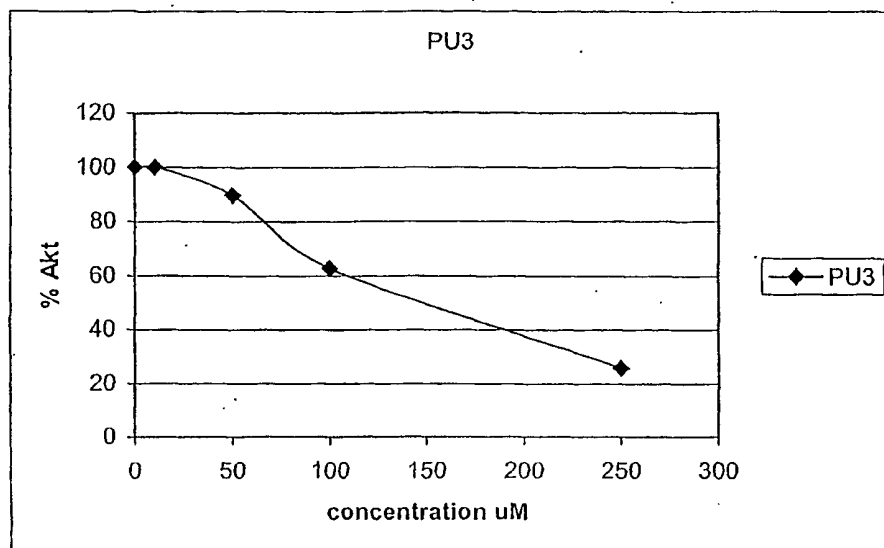


Fig. 17A

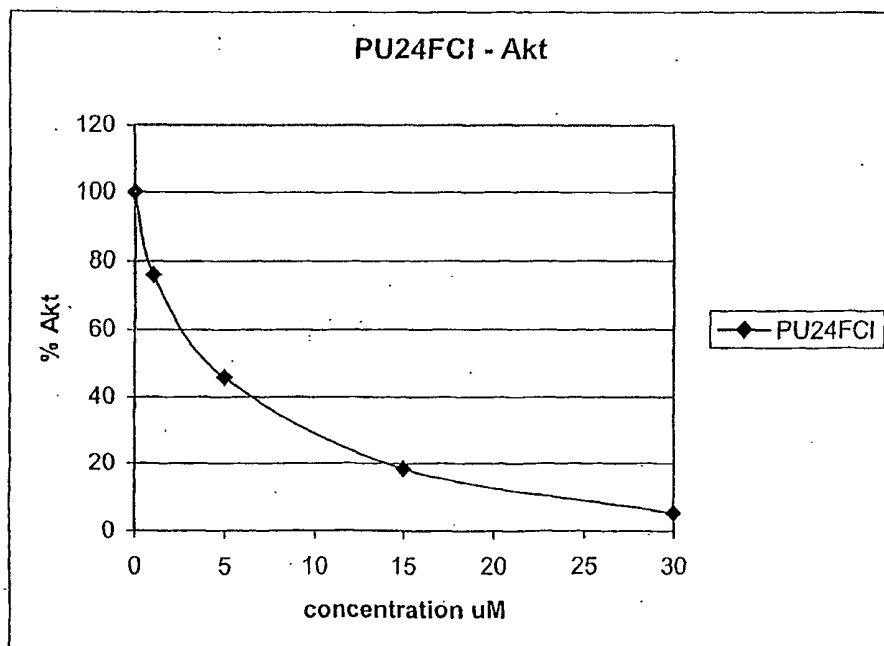


Fig. 17B

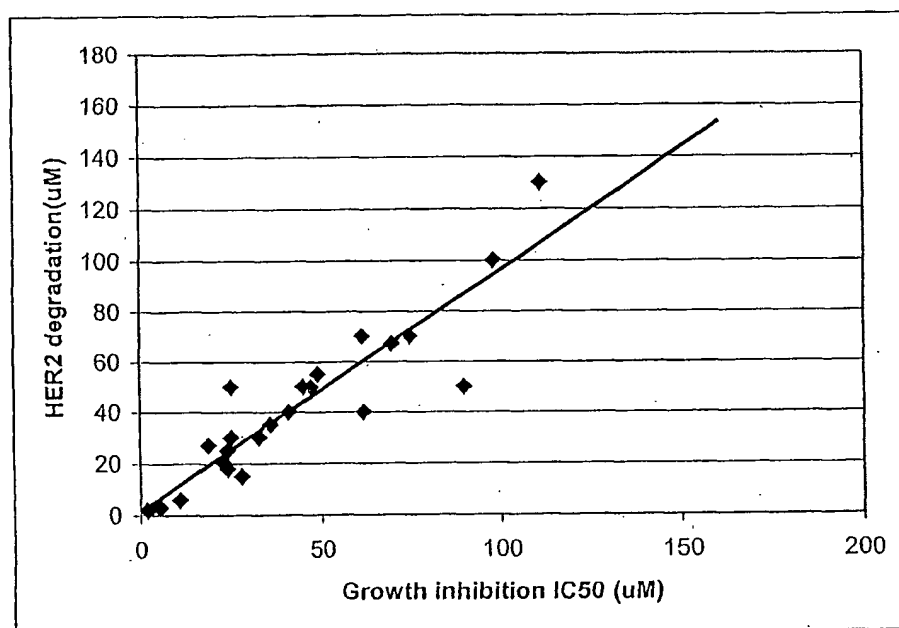


Fig. 18A

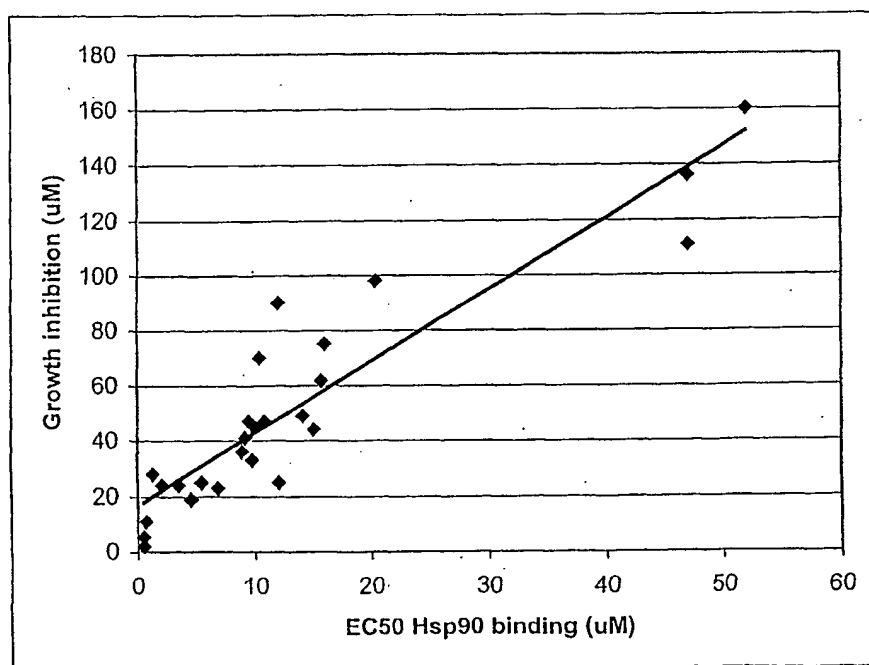
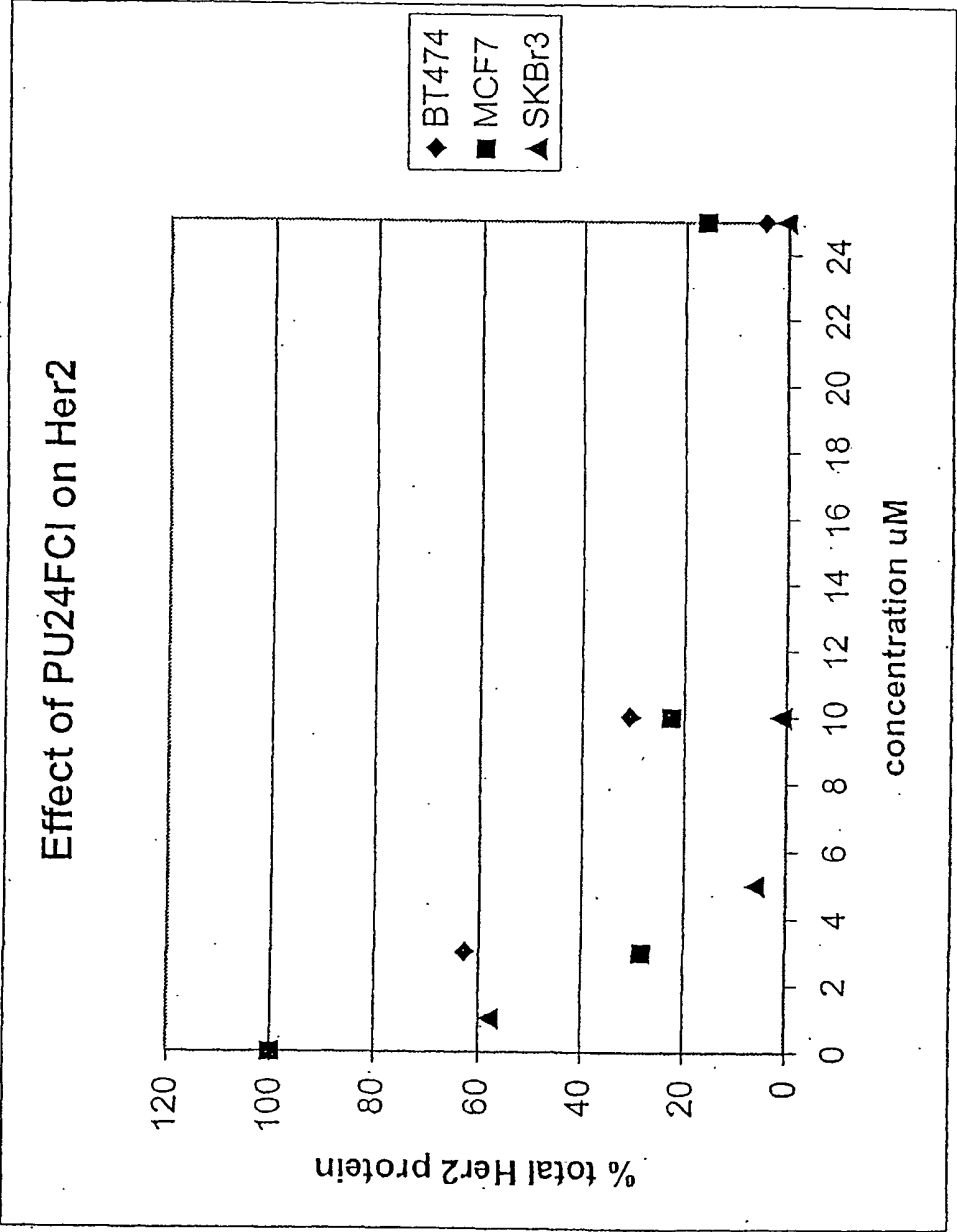


Fig. 18B

Fig 19



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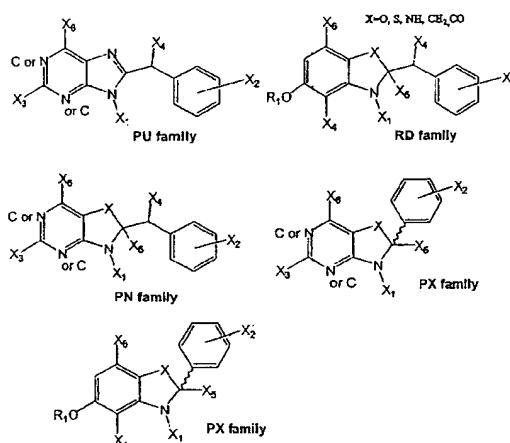
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[Continued on next page]

(54) Title: SMALL MOLECULE COMPOSITIONS FOR BINDING TO HSP90



(57) Abstract: Structural differences in binding pockets of members of the HSP90 family are exploited to achieve differential degradation of kinases and other signaling proteins by using small molecules which interact with the N-terminal binding pocket with an affinity greater than ADP and different from ansamycin antibiotics for at least one HSP90 family species (Figure 5). These small molecules are soluble in aqueous media, providing a further advantage over ansamycins. Pharmaceutical compositions contain a carrier and a molecule that includes a moiety which binds to the N-terminal pocket of at least one member of the HSP90 family. Such binding moieties have anti proliferative activity against tumor cells which depend on proteins requiring chaperones of the HSP90 family. Different chemical species have different activity, allowing the selection of Her2 degradation without degradation of Raf kinase. Thus, the binding moieties possess inherent targeting capacity. The small molecules can be linked to targeting moieties to target the activity to specific classes of cells. The invention includes treatment of diseases, including cancers. Dimeric forms of the binding moieties may be employed.



CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

(88) Date of publication of the international search report:

5 June 2003

Published:

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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/46303

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C07D 487/04, 498/04, 513/04, 471/04, 209/36, 209/08, 209/12, 209/18, 209/22, 263/56, 263/57, 235/26, 235/08, 235/12, 235/16, 235/18, 277/64, 277/66, 473/00, 473/02, 473/22, 473/18, 473/24, 473/28, 473/30, 473/34, 473/36, 473/40; A61K 31/519, 31/437, 31/404, 31/423, 31/4184, 31/428, 31/ 52, 31/522, GOIN 33/53, 33/543; A61P 35/00

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
U.S. : Please See Continuation Sheet

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
CAS ONLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 6,413,975 B1 (CHASIN et al) 2 July 2002 (02.07.2002). See Example 4.	18
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A		19-20
A	US 6,440,982 B1 (MAW et al) 27 August 2002 (27.08.2002). See entire document	6-9, 17
A	BLACK et al. "Reaction Of Ninhydrin With Activated Anilines: Formation Of Indole Derivatives." Tetrahedron. 1994, Vol. 50, No. 37, pages 10983-10994. See Compound 17.	1-5, 10-16
A	KATO et al. "Synthesis Of Compounds Related To Antitumor Agents IV. On The Reaction Of Aromatic Carboxylates With 2,4 Diamino-5-hydroxy-6-methylpyrimidine." Chem. Pharm. Bull. 1976, Vol. 24, No. 10, pages 2461-2469. See Table 2.	1-5, 10-16
A	US 6,210,974 B1 (GOLD) 3 April 2001 (03.04.2001). See Claims 1-2.	21-26
Y,E	US 6,335,157 B1 (GONZALEZ et al) 1 January 2002 (01.01.2002). See Claims 7-18.	21-26

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"B" earlier application or patent published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

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Authorized officer

John D. Robert for
MARK BERCH

Telephone No. 703-308-1235

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/46303

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claim Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claim Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claim Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:
Please See Continuation Sheet

1. ☒ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐

The additional search fees were accompanied by the applicant's protest.

☒

No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

PCT/US01/46303

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING

This ISA found multiple inventions as follows:

Group I, claims 1,2 (part), 4-5 (part), 10-12 (part), 13, 14 (part), 15, 16 (part), 17-20, drawn to purines.

Group II, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 16 (part), drawn to pyrrolopyrimidines.

Group III, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to oxazolopyrimidines.

Group IV, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to thiazolopyrimidines.

Group V, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to Imidazopyridines.

Group VI, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to pyrolopyridines.

Group VII, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to oxazolopyridines.

Group VIII, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to thiazolopyridines.

Group IX, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to Indoles.

Group X, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to Benzoxazoles.

Group XI, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to Benzimidazoles.

Group XII, claims 1, 2 (part), 3, 4-5 (part), 6-9, 10-12 (part), 14 (part), 16 (part), drawn to Benzothiazoles.

Group XIII, claims 21-26, drawn to screening methods.

Claims 1, 2, 4 and 11 do not recite any structure and hence the full scope cannot be determined. These claims are searched only to the extent that they read on the elected invention(s).

The inventions listed as Groups I-XIII do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

Group I-XII each have the distinct special technical feature of the particular binding moiety. Groups I-IV are based on fused pyrimidines. In each case, the pyrimidine is fused to a different heterocyclic ring, viz, imidazole, pyrrole, oxazole and thiazole. These four rings differ in the nature of the heterocycle present, respectively, N2C3, NC4, ONC3 and SNC3, Groups V-VIII are fused pyridines, fused in each case with the imidazole, pyrrole, oxazole and thiazole rings. Groups IX-XII have cores with only one heterocyclic ring, in which the 4 rings are each fused to a benzene ring.

Group XIII has the special technical feature of performing a measurement. Note that Group XIII does not constitute a method of use of any of the compounds of Groups I-XII because a) there is no actual use of a compound, only on determination of its property and b) Group XIII does not require the use of any compound of employed in any of claims 1-17, since the method of Group XIII does not require that compound falling within claim 1 be employed.

INTERNATIONAL SEARCH REPORT

PCT/US01/46303

Continuation of B. FIELDS SEARCHED Item 1:

544/264, 265, 267, 271-273, 276-278, 280; 546/113-116, 183; 548/152, 165, 169, 170, 178-180, 217, 224, 306.4, 309.4, 310.7, 304.4, 484, 490, 491; 514/260.1, 263.3, 263.34, 263.37, 263.38, 263.4, 300-303, 265.1, 367, 375, 387, 394, 415, 418, 419; 436/501, 503